

Cite this: *Dalton Trans.*, 2019, **48**, 4894

Applications of boroxide ligands in supporting small molecule activation by U(III) and U(IV) complexes†

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The boroxide ligand [OBAr₂][−] (Ar = Mes, Trip) is shown to be able to support both U^{III} and U^{IV} centres for the first time. The synthesis and structures of homoleptic and heteroleptic U^{III} and U^{IV} complexes are reported. The UX₃ complex with larger substituents, [U(OBTrip)₂]₂, exhibits greater thermal stability compared to less encumbered [U(OBMes)₂]₂ but reacts with a smaller range of the small molecules tested to date. Initial studies on their capacity to participate in small molecule chemistry show that dark purple [U(OBMes)₂]₂ binds and/or reductively activates a variety of small molecules such as pyridine-oxide, triphenylphosphineoxide, sulfur, and dicyclohexylcarbodiimide. While [U(OBMes)₂]₂ shows no reaction with CO or CO₂, [U(OBTrip)₂]₂ is oxidised by both, in the former case forming [U(OBTrip)₂]₄, and in the latter case forming a small quantity of the structurally characterised μ -carbonate product [(μ -CO₃){U(OBTrip)₂}]₂.

Received 21st December 2018,

Accepted 28th February 2019

DOI: 10.1039/c8dt05051a

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Introduction

Complexes of the form UX₃ where X is a large, monoanionic ligand such as an amide or aryloxy have shown great utility in the binding and reductive activation of small molecules. The choice of X provides great control over the space at the Lewis acidic metal, and the strength of the reducing capacity of the U^{III} centre. For example, the common silylamide and aryloxy complexes [UN^{''}]₃ and [U(ODtbp)₃] (N^{''} = N(SiMe₃)₂, ODtbp = O-2,6-^tBu₂C₆H₃) reductively couple carbon monoxide gas at room temperature and pressure to form the [OCCO]^{2−} fragment as the sole product in [X₂UOCCOUX₂].^{1,2} The dinuclear tren derivative [{U(tren')}]₂(μ - η^2 : η^2 -N₂) (tren' = N(NCH₂CH₂NSi^tBuMe₂)₃) and [U(ODtbp)₃] show ready but reversible side-on N₂ coordination,² but the OTtbp analogue (OTtbp = O-2,4,6-^tBu₃C₆H₂) [{U(OTtbp)₃}]₂(μ - η^2 : η^2 -N₂) releases

N₂ only upon heating a solution to 80 °C, even in the presence of small O-containing reagents such as CO₂ and CO.²

We showed that bulky trimesitylsiloxide complexes bind N₂ even more strongly in [{U(OSiMe₃)₃}]₂(μ - η^2 : η^2 -N₂),³ and others have shown that further functionalisation of the trapped and reduced N₂ in polynuclear nitrido complexes supported by *tert*butylsiloxides is possible.⁴ Flexible uranium siloxides have also allowed reactions such as CO₂ deoxygenation and carbonate formation in [{U(OSi(O^tBu)₃)₃}]₂(μ - η^1 : η^2 -CO₃).⁵

The capacity to transfer one X ligand has also provided significant advantages for new reactivity. For example, storage of arene solutions of [UX₃] (X = N^{''}, ODtbp) results in the spontaneous reductive activation of the arene (over 90 h at 90 °C, or 30 days at 25 °C) through the formation of [X₂U]₂(μ - η^6 : η^6 -C₆H₆) and elimination of two equivalents of UX₄ by-product.⁶ The trapped arenes are also sufficiently reduced that a mild, homogeneous borylation of one C–H group is possible.

Boroxides provide an interesting alternative to alkoxides and aryloxides as these can provide greater delocalisation of the electron density across the OBM bond, and even a distant Lewis acidity via an empty orbital on the boron atom, thus allowing the electron density to be tuned at the metal centre.⁷ The aryl functionalisation can also provide a larger protected pocket at the metal as the bulk of the aryl groups is maintained at a greater distance from the metal binding site.⁸

We have studied the capacity of the boroxide ligand [OBAr₂][−] (Ar = Mes, Trip) to bind U^{III} for the first time. Here we show the synthesis and structures of new U^{III} and U^{IV} com-

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† Electronic supplementary information (ESI) available: General experimental considerations, preparation of reagents and additional experimental details. CCDC 1886802–1886811, 1886182 and 1886183 (complex 11^t and 2^t). For ESI and crystallographic data in CIF or other electronic format see DOI: 10.1039/c8dt05051a

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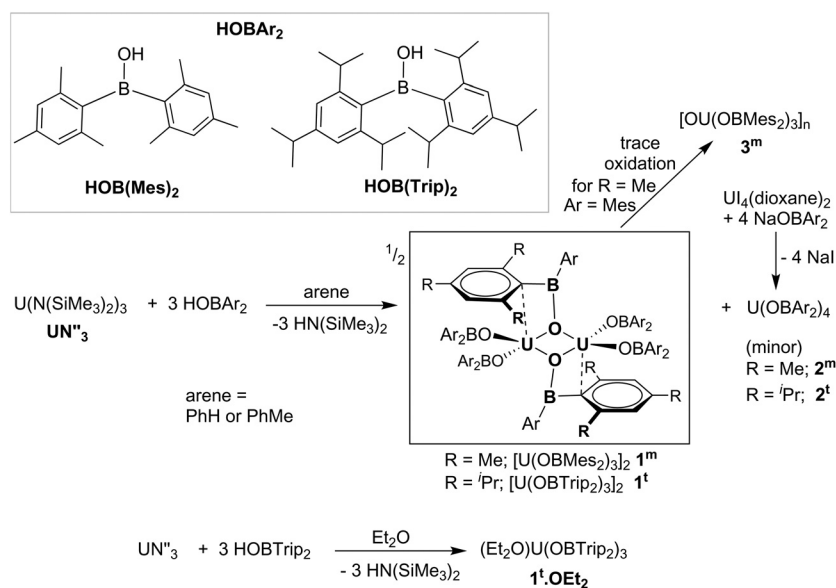
plexes and initial studies on their capacity to participate in small molecule chemistry.

Results and discussion

Syntheses

The formation of dark purple $[U(OBMe_2)_3]_2$ **1^m** from $HOBMe_2$ and UN''_3 occurs in less than 3 hours at room temperature in toluene solution, Scheme 1. Work-up from hexanes affords dark crystals in 46% yield, suitable for a single crystal X-ray diffraction study, Fig. 1. At room temperature the 1H NMR spectrum of **1** shows the retention of the dimeric structure in solution with two different sets of ligand resonances assignable to terminal and bridging ligands. The bulkier aryl groups on the boroxide still lead to similar reaction chemistry to afford $[U(OBTrip_2)_3]_2$ **1^t**, although the major by-product in

the synthesis is the tetrakis(ligand) complex **2^t**, and both UX_3 and UX_4 are isolated by fractional crystallisation in similar (*ca.* 30%) yields. The disproportionation of uranium(III) to uranium(IV) and uranium metal is a known oxidative pathway.^{6,9,10} Dark red crystals of the mononuclear diethylether adduct $[(Et_2O)U(OBTrip_2)_3]$ **1^t-Et₂O** are formed from reactions in diethyl ether and have also been structurally characterised by a single crystal X-ray diffraction study, Fig. 1. The 1H NMR spectrum of **1^m** suggests exchange between unbound and bound Ar groups on bridging $OBMe_2$ at ambient temperature and support the dimeric conformation. The resonances assigned to the bridging ligands are strongly paramagnetically shifted out of the diamagnetic region (from -3.71 to -15.13 ppm) due to proximity to the uranium centres, whereas the resonances corresponding to the terminal ligands (double the relative intensity) are between 8.29 to 2.53 ppm. The ^{11}B NMR spectrum contains a resonance at 73.7 ppm that is tentatively assigned to one of



Scheme 1 Synthesis of complexes **1** showing common by-products and decomposition products.

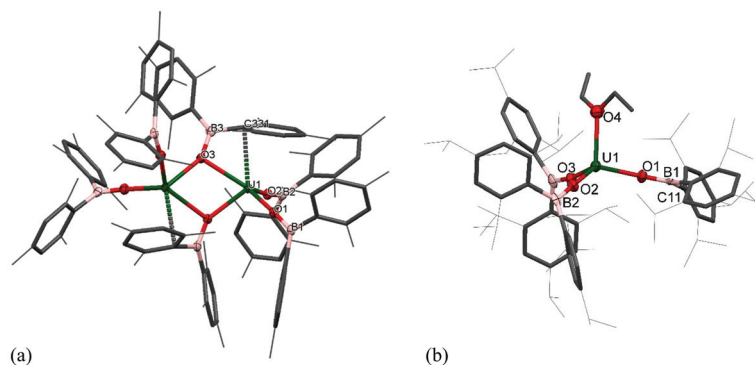
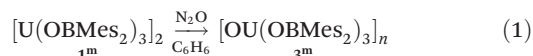


Fig. 1 Displacement ellipsoid drawing of the molecular structure of (a) **1^m** and (b) **1^t-Et₂O**. The hydrogen atoms are omitted and the mesityl and triisopropylphenyl groups are depicted as wireframe and capped sticks for clarity. The thermal ellipsoids are displayed at 50% probability.



what should be two different boron environments, but only one other signal attributed to trace free ligand (50.8 ppm) is observed over an extended frequency range.

Clean starting materials and precise control of the stoichiometry is key to the formation of pure samples, as both derivatives of **1** are very air sensitive. The homoleptic U^{IV} complexes $[U(OBAR_2)_4]$ **2** are significantly more stable, and can be formed as minor side products in syntheses of **1**, and from independent metathetical routes from the reaction between uranium tetraiodide and the salt $NaOBAr_2$, Scheme 1. A high-yielding route to **2^m** is available from the reaction of $NaOBMe_2$ with $[UI_4(1,4\text{-dioxane})_2]$ in toluene, affording $[U(OBMe_2)_4(\text{dioxane})_{0.5}]_2$ (**2^m-diox**). Colourless crystals of **2^m-diox** and **2^t-C₇H₈** suitable for XRD analysis were obtained from slow diffusion of hexanes into the reaction mixture, and standing of a concentrated toluene solution, respectively, see below. Complex **2^m-diox** was characterised by 1H NMR spectroscopy, mass spectrometry (APPI-MS) and single crystal XRD analysis. The 1H NMR spectrum shows the corresponding signals of one boroxide ligand environment. Mass spectrometric analysis showed the molecular ion at $m/z = 1298.76$ that corresponds to $[M]^{++}$ fragment $[U(OBMe_2)_4]^{++}$. The 1H NMR spectrum of **2^t** contains many more resonances than that of **1^t** due to the congestion around metal centre; twelve magnetically different methyl group resonances are measured in the range of 4.13–18.40 ppm and five different methine groups are discernible in the range 8.86–4.78 ppm.



Additionally, complex **1^m** is readily oxidised to the oxo U^{IV} product $[OU(OBMe_2)_3]_n$ **3^m**, due to the presence of traces of oxygen, which has been fully characterised from an independent synthesis. Samples also decompose when heated to 100 °C in toluene or in the solid state (10^{-3} mbar, 115 °C, 6 h). Upon exposure of a benzene solution of complex **1^m** to one atmosphere of dry N_2O , a rapid colour change from dark purple to light brown is observed due to the formation of the U^{IV} oxo complex **3^m**, which is assigned as containing bridging oxo ligands by comparison with the literature and in consideration of the size of the $OBMe_2$ ligands, eqn (1). The 1H and ^{13}C NMR spectra show resonances corresponding to one mesityl environment, which is slightly shifted due to paramagnetic contributions. The ^{11}B NMR spectrum contains one resonance at 89.29 ppm, which confirms the presence of only one boron environment. The elemental analysis result is in agreement with the formation of **3^m**. Reactions designed to reduce complex **3^m** on a bulk scale have so far been unsuccessful; a small quantity of a U^{III} product $K_2[\{(OBMe_2)_3U\}_2(\mu-O)_2]$ (**10^m**) that was isolated from a reaction with KC_8 has been structurally characterised (see ESI† for details).

Any traces of the metallacycle $[UN''_2\{\kappa^2-N'(SiMe_2CH_2)\}]$ ($N' = N(SiMe_3)$, a common U^{IV} contaminant in UN''_3 , forms $[UN''_3(OBMe_2)]$ and $[UN''_3(OBTrip_2)]$ (*vide infra*).

Crystallography

The solid-state structure of **1^m** confirms the formation of a centrosymmetric binuclear U^{III} complex in which the two uranium centres are bridged by two boroxide ligands (Fig. 1a) in a distorted trigonal bipyramidal geometry due to the interaction of the uranium centres with the bridging mesityl groups. The $U\cdots C$ distances for these interactions ($U\cdots C_{avg}$ 2.845 Å) are slightly longer than other η^1 interactions with uranium, such as the complexes bearing the ligand $\{1,3[2,5-(^iPr)_2PhNC(=CH_2)]C_6H_4\}$ reported by Gambarotta and co-workers (2.678(12) Å).¹¹ The $U-O$ bond lengths for the boroxide ligands are 2.548(10) Å and 2.372(10) Å for the bridging boroxides, and an average $U-O_{avg}$ bond length of 2.196 Å for the terminal boroxides. Both are in the same range as the $U-O$ distances for the siloxide-bridged U^{III} dimer reported by Mazzanti and co-workers, $[U(OSi(O^tBu)_3)_2(\mu-OSi(O^tBu)_3)]_2$, which features two U^{III} centres with two bridging and two terminal siloxide ligands with $U-O$ distances of 2.549(3) Å and 2.396(3) Å for the bridging siloxides, and a $U-O_{avg}$ of 2.193(4) Å for the terminal ligands. Moreover, the $U\cdots U$ distance for complex **1^m** is 3.966 Å, which is similar to the U^{III} siloxide complex ($U\cdots U$ 3.9862(2) Å).⁵ The $U-O_{avg}$ distance for the terminal ligands is similar to that of $[(Et_2O)U(OBTrip_2)_3]$ **1^t-Et₂O** (mean 2.183(7) Å) and slightly longer than that of the uranium tris(aryloxide) complex, $[U(ODtbp)_3]$, with $U-O_{avg} = 2.159$ Å. The average $B-O$ bond length of complex **1^m** is 1.351 Å. For both complexes the $B-O_{avg}$ fall within range for Ar_2BO-M complexes reported in the literature (1.295–1.405 Å, mean 1.350(1) Å for a fragment search for Mes_2BO-M in the Cambridge Structural Database).¹² In the solid-state structure of **1^t-Et₂O** the average $U-O_{boroxide}$ distance is 2.183 Å, slightly shorter in **1^m** but significantly longer than in $[U(ODtbp)_3]$ (2.159 Å). The $U-O_{ether}$ is 2.530(7) Å and is much shorter than in the seven-coordinate complex $[(^{Neop}ArO)_3tacn]U(OEt_2)$ (2.669(2) Å, $^{Neop}ArO = 2\text{-neopentyl-4-methylphenoxide}$, $tacn = \text{triazacyclononane}$), which is the only other crystallographically characterised U^{III} diethyletherate.¹³ This is consistent with both the more electron-deficient uranium centre engendered by the poor π donor properties of the boroxide ligand and the lower coordination number in **1^t-Et₂O**.

Together, the nearly linear $U-O-B$ angle for the terminal boroxide ligands (mean 171.6° in **1^m** 171.0° in **1^t-Et₂O**) and the elongation of the $U-O$ bonds are a direct consequence of the reduced π -donor ability of the ligand compared with a carbocyclic aryloxide, which is decreased by having a boron substituent on the ligand. Electron density from the oxygen is accepted by the boron, leading to a lower degree of π -donation from the boroxide to the uranium centre.

The solid-state structure of **2^m-diox** reveals two U^{IV} ions both in a distorted trigonal bipyramidal coordination environment (Fig. 2a). Each uranium centre is ligated by one end of a bridging dioxane molecule in an axial position with a $U-O$ distance of 2.577(3) Å, which is considerably longer than in $[UI_4(1,4\text{-dioxane})_2]$ or the $[UI_2(\text{dioxane})_2(\text{aryloxide})]$ previously reported by our group (mean = 2.333(6) Å and 2.087 Å respectively).^{14–16} The



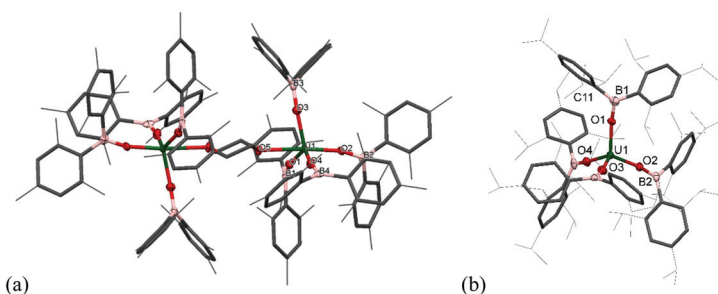


Fig. 2 Displacement ellipsoid drawing of the molecular structure of (a) $2^m \cdot \text{diox}$ and (b) $2^t \cdot \text{C}_7\text{H}_8$. Thermal ellipsoids are displayed at 50% probability. Lattice solvent and hydrogen atoms omitted, and aryl substituents are depicted as wireframe and capped sticks for clarity.

boroxide ligands in the equatorial positions have a $\text{U}-\text{O}_{\text{avg}}$ distance of $2.144(3) \text{ \AA}$, which is shorter than in the uranium(III) complex 1^m , consistent with a U^{IV} centre, and similar to the $\text{U}-\text{O}$ bond length for $[\text{U}(\text{OBTrip})_4]$ with a $\text{U}-\text{O}$ distance of $2.159(5) \text{ \AA}$. The boroxide ligands in the axial positions have a $\text{U}-\text{O}$ distance of $2.095(3) \text{ \AA}$, which is much shorter than the equatorial boroxides. The $\text{B}-\text{O}_{\text{avg}}$ distance is $1.356 \text{ \AA}$, which is similar to complex 1^m and falls within range for reported metal boroxide complexes. The $\text{U}-\text{O}-\text{B}_{\text{avg}}$ angles are 167.1° , which is comparable to the $\text{U}-\text{O}-\text{B}$ angles of the terminal boroxide ligands in 1^m (mean 171.6°).

Single crystals of $2^t \cdot \text{C}_7\text{H}_8$ suitable for X-ray diffraction analysis were obtained from concentrated toluene solution, and its solid-state structure is depicted in Fig. 2b. The solid-state structure shows an unsolvated four-coordinate uranium centre in tetrahedral geometry. The mean boroxide ligand $\text{U}-\text{O}$ distance of $2.159(5) \text{ \AA}$ is shorter than in the uranium(III) complex $1^t \cdot \text{Et}_2\text{O}$, consistent with a uranium(IV) centre. The $\text{U}-\text{O}$ bond lengths in 2^t are longer than those in homoleptic uranium(IV) aryloxide $[\text{U}(\text{ODtbp})_4]$ ($2.135(4) \text{ \AA}$), due to the greater steric demand of the boroxide ligand and poorer metal to ligand backbonding. The $\text{U}-\text{O}-\text{B}$ angles are comparable to those in $1^t \cdot \text{Et}_2\text{O}$ (mean $171.0(5)^\circ$), despite the considerable bulk about the uranium centre. The average $\text{B}-\text{O}$ bond length of $1.364(7) \text{ \AA}$ falls within range for reported metal boroxide complexes and is slightly longer than in $1^t \cdot \text{Et}_2\text{O}$. The longer $\text{B}-\text{O}$ bond may be due to the more Lewis acidic uranium(IV) centre in 2^t .

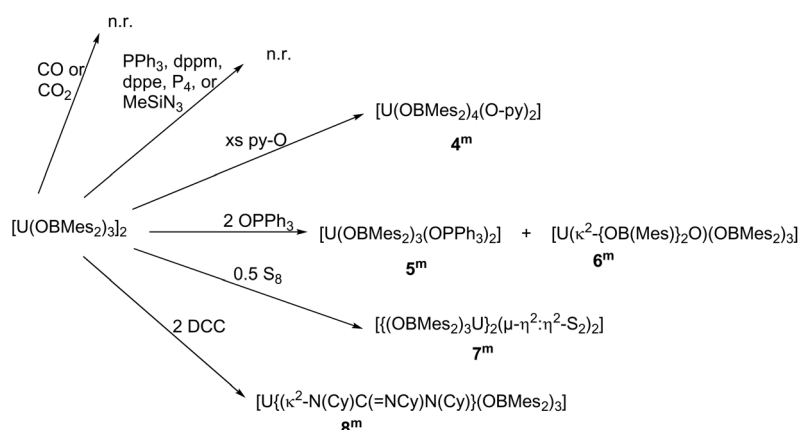
Reactions of 1^m and 1^t

Compound 1^m has been treated with a wide range of small molecules, Scheme 2.

Reactions with small, neutral N- or P-donor reagents that are readily visible by NMR spectroscopy were designed to identify whether the boron atoms in the ligands retain sufficient Lewis acidic character to participate in reactions. However, arene solutions of 1^m showed no reaction with added Me_3SiN_3 , PPh_3 , $\text{CH}_2(\text{PPh}_2)_2$ (dppm), $\text{C}_2\text{H}_4(\text{PPh}_2)_2$ (dppe) or white phosphorus, P_4 . Normal, simple UX_3 complexes with large monoanionic X ligands would be anticipated to react with these, readily forming oxidised uranium complexes or adducts, for example, $[\text{Me}_3\text{SiN}=\text{UN}''_3]$,^{17,18} $[\text{Me}_3\text{SiN}=\text{U}\{\text{OSi}(\text{O}^t\text{Bu})_3\}_3]$,¹⁹ $[\text{U}(\text{BH}_4)_3(\text{PET}_3)_2]$.²⁰

Pyridine N-oxide

The formation of oxo complexes with structures such as $\text{UL}_n=\text{O}$,^{21–23} $\text{L}_n\text{U}\equiv\text{O}$ ^{13,24} or $\text{U}-\text{O}-\text{U}$ ²⁵ have been reported from reactions with pyridine N-oxide. In this case, addition of benzene to a mixture of $[\text{U}(\text{OBMes}_2)_3]_2$ and an excess of pyridine N-oxide provides a brown solution from which $[\text{U}(\text{OBMes}_2)_4(\text{py}-\text{O})_2]$ (4^m) can be isolated as dark orange solids after 72 hours, Scheme 2. The reaction proceeds through oxidation of complex 1^m to mononuclear U^{IV} complex 4^m with ligand redistribution. Coordination of pyridine



Scheme 2 Reactions of complex 1^m investigated.

N-oxide to a metal is not unique, although relatively rare.^{13,26–29} The identity of complex **4^m** was confirmed by ¹H NMR spectroscopy, APPI mass spectrometry and single crystal XRD analysis. Mass spectrometric data showed the expected molecular ion at *m/z* = 1223.65, which corresponds to the [M]⁺⁺ fragment, [U(OBMes₂)₄(Py–O)₂]⁺⁺. Dark orange crystals of **4^m** suitable for XRD analysis were grown from a concentrated benzene solution. The data are of sufficient quality to discuss connectivity information but not for bond metrics. The solid-state structure reveals a U^{IV} complex with a distorted octahedral geometry with the boroxide ligands in the equatorial positions and the pyridine *N*-oxide ligands in the axial positions (Fig. 3).

Triphenylphosphine oxide

Phosphine oxides are widely used ligands in f-element chemistry. As a result of the oxophilic and Lewis acidic character of lanthanides and actinides, phosphine oxides coordinate readily, and are able to stabilise a variety of complexes in low and high oxidation states.^{30–35} Moreover, applications in catalysis and liquid–liquid extraction processes have been found.³⁶ Therefore, the reaction of a benzene solution of the uranium boroxide complex **1^m** with two equivalents of OPPh₃ was carried out, Scheme 3. This leads to the formation of [U(OBMes₂)₃(OPPh₃)₂] (**5^m**), which co-crystallised together with a minor by-product [U(κ²-{OB(Mes)}₂O)(OBMes₂)₃] (**6^m**).⁷

The ¹H NMR spectrum of **5^m** in C₆D₆ contains six resonances which range from 6.9 to 1.87 ppm. The ³¹P NMR spec-

trum shows a single broad resonance at –17.83 ppm, and this is significantly shifted from the free OPPh₃ resonance (~25 ppm) as a result of its proximity to the metal centre. Mass spectrometric analysis (APPI-MS) showed a molecular ion peak at *m/z* = 1589.77 that corresponds to fragment [5^m]⁺. By-product [U(κ²-{OB(Mes)}₂O)(OBMes₂)₃] (**6^m**) contains a co-ordinated boroxinato ligand, a cyclic anhydride of borinic acid,³⁷ which is formally a product of ligand condensation, but probably formed from a contaminant in the ligand. Boroxines are rigid bidentate ligands which have drawn attention recently, mostly by forming complexes of main group elements, e.g. for Al;^{38–40} Sn, Sb and Bi;^{41–47} but also for transition metals such as Mn,⁴⁸ Au^{37,49} and Pt.⁵⁰

Single crystals of **5^m** and **6^m** were obtained by slow diffusion of hexanes into a benzene solution. Complex **5^m** has an approximately trigonal bipyramidal coordination geometry (Fig. 4). The U–OPPh₃ bond lengths are 2.273(4) Å and 2.270(4) Å, which are ~0.1 Å shorter than other U^{III}–OPPh₃ distances previously reported.^{36,51} The U–O–P angles for the phosphine oxide ligands are 170.4(3)° and 174.4(2)° respectively, and the O₄–U–O₅ angle is 176.62(13)°. These angles are similar to the angles in the known complex, [U(NPhAr^F)₃(OPPh₃)₂].²⁴ The U–O_{avg} distance for the boroxide ligands is 2.124 Å, which is ~0.07 Å shorter than the U–O bond lengths for the terminal boroxides in **1^m** (mean 2.196 Å). Similarly, the average B–O distance is 1.345 Å which is slightly shorter than complex **1^m** (mean 1.351 Å). Moreover, the mean U–O–B angle is 176.0°, and this is less bent than the terminal boroxide in **1^m** (mean 171.6°). Complex **6^m** has a nearly square pyramidal coordination geometry. The metallacycle formed from the [OBOBO]^{2–} unit has similarities with early metal complexes such as (η²-catechol-μ-catecholborate)₃ThCl(C₄H₈O)₃·C₄H₈O that are coordinated by alternating catechol and catecholborate groups in a hexaoxo trianionic macrocyclic structure, observed for Zr, Nd, Sm, U and Th centres.^{52,53} For the boroxine ligand, the U–O bond distances are 2.193(4) Å and 2.191(4) Å. The B–O distances range from 1.342(9)–1.370(9) Å, which are in the same range as previously reported for main group boroxines. The six-membered ring unit has internal angles close to 120°, except for the O₉–U–O₁₁ bond which is much smaller, being only 77.40(16)°. The boroxide ligands have similar U–O and B–O bond lengths (U–O_{avg} 2.172 Å, B–O_{avg} 1.316 Å) to the terminal boroxides in **1^m** (mean U–O 2.196 Å,

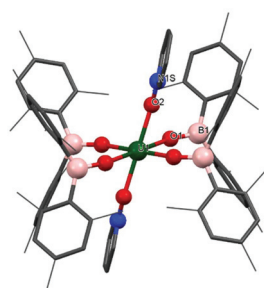
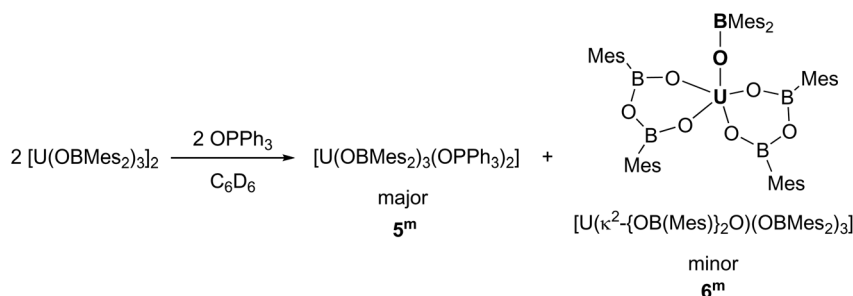


Fig. 3 Solid-state structure of **4^m**. Hydrogens atoms are omitted and mesityl and pyridine oxide ligands are depicted as wireframe and capped sticks for clarity.



Scheme 3 Synthesis of compounds **5^m** and **6^m**.



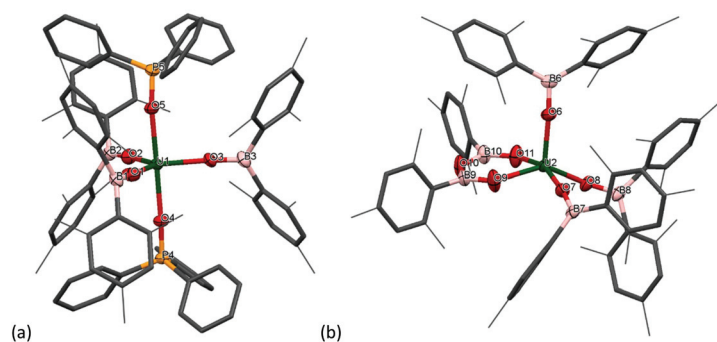


Fig. 4 Solid-state structure of (a) **5^m** and (b) **6^m**. Mesityl and phenyl ligands are depicted as capped sticks and wireframe for clarity. Hydrogen atoms are omitted. The thermal ellipsoids are displayed at 50% probability.

B–O 1.351 Å). Finally, the U–O–B angles for the boroxides range from 155.2(4)° to 168.7(4)°, which are significantly more bent than for the terminal boroxides in **1^m** (mean 171.6°).

Elemental sulfur

The reactivity of chalcogens and chalcogenides with actinide complexes has gained attention recently. Of particular interest is the potential of the metal–ligand bonding in these more polarisable systems to advance our understanding of the degree of covalency in uranium–ligand bonds, as this can provide insights into better separation of actinides from lanthanides in spent nuclear fuel.^{54–58} Addition of 0.5 equivalents of S₈ to a stirring solution of [U(OBMeS₂)₃]**1^m** leads to the formation of [{(OBMeS₂)₃U}₂(μ-η²:η²-S₂)₂] (**7^m**), which contains two independent bridging S₂ units. There is only one other example in the literature of a uranium complex featuring two bridging S₂ units, [({(AdArO)₃N}U)₂(μ-η²:η²-S₂)₂].⁵⁹ The ¹H NMR spectrum contains a set of resonances corresponding to the OBMeS₂ ligand, indicating one ligand environment. Mass spectrometric analysis shows the molecular ion at *m/z* = 1033.56 that corresponds to the [M]⁺ fragment, [{(OBMeS₂)₃U}(μ-η²:η²-S₂)₂]⁺. Translucent red crystals of the sulfur-bridged complex **7^m** suitable for XRD analysis were grown from a concentrated toluene solution.

Fig. 5 shows the molecular structure of **7^m**, confirming the formation of a uranium dimer with two S₂ units. Each uranium ion has a distorted pentagonal bipyramidal coordination geometry. The S_{1B}–S_{2B} bond length is 2.050(2) Å, which is a typical value for a S–S single bond, and comparable to H₂S₂ with a S–S bond length of 2.055 Å.⁴⁴ The S...S distance between the two units is 3.491 Å, excluding the possibility of a S₄^{2–} unit.

Thus, although one would normally anticipate a U^{IV} formal oxidation state product to form from the reduction of sulfur, here the U^V assignment appears more appropriate. The ligands are both identified as bridging persulfido (S₂^{2–}), giving a formal U^V oxidation state for **7^m** by comparison of the U–E bonds. Previously reported uranium persulfido S₂^{2–} complexes show S–S distances from 2.050 Å to 2.103 Å.^{60–68} Previously reported transition metal supersulfido S₂[–] complexes show S–S

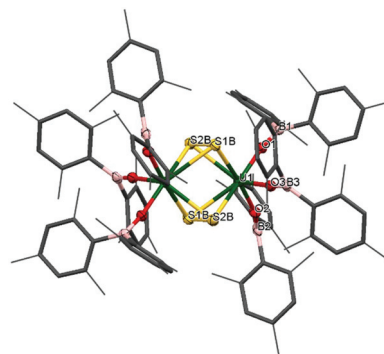


Fig. 5 Solid-state structure of complex **7^m**. Mesityl ligands are depicted as capped sticks and wireframe for clarity. Hydrogen atoms are omitted. The thermal ellipsoids are displayed at 50% probability.

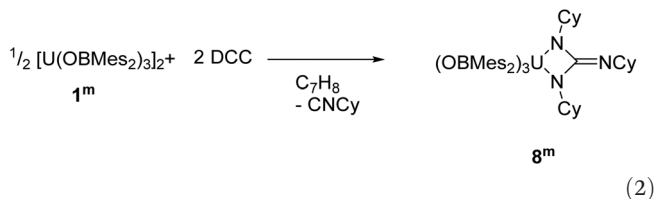
distances from 1.944 to 2.023 Å.^{69–72} Furthermore, the U–O_{avg} bond length is 2.078 Å, which is significantly shorter than the U–O bond lengths for the terminal boroxides in U^{III} **1^m** (mean 2.196 Å), and in **2^m** (mean 2.119 Å), also suggesting a formal oxidation state of +5 is most likely. Moreover, the B–O average distance is 1.359 Å, which is slightly longer than for complexes **1^m** and **2^m** (mean 1.351 Å and 1.356 Å respectively). The U–O–B angle of 165(3)° is much more bent than for the terminal boroxide ligands in **1^m** (mean 171.6°), which is reasonable if more electron density from the O atoms are being distributed into a backdonation from the uranium centres to the sulfido ligands.

DCC, *N,N'*-dicyclohexylcarbodiimide

The direct reaction of carbodiimides such as (DCC = *N,N'*-dicyclohexylcarbodiimide) with uranium was first reported in 2009.⁷³ The aim of these reactions was to insert a carbodiimide into a U–C ligand bond in order to increase the steric crowding in organouranium complexes. Since then, several examples of insertion of carbodiimides into metal–ligand σ-bonds have been reported,^{74–77} as well as further reactivity towards small molecules with these ligands.^{78–80} Moreover, reactions of carbodiimides have also led to the formation of metallacycles that



have unique structural properties and possible applications in catalysis.^{81–84}



Dropwise addition of a toluene solution of DCC to a dark purple toluene solution of **1^m** produced a green solution from which $[\text{U}\{\eta^2\text{-N}(\text{Cy})\text{C}(\text{=NCy})\text{N}(\text{Cy})\}(\text{OBMe}_2)_3]$ (**8^m**, Cy = *cyclo*-C₆H₁₁), eqn (2), was isolated in close to quantitative yield after work-up. Mass spectroscopic analysis confirms the formulation of **8^m** as the molecular ion at $m/z = 1337.85$ that corresponds to $[\text{M} + \text{H}]^+$ was observed. Single crystals for XRD characterisation could not be obtained. The ¹H NMR spectrum of **8^m** has resonances from 58.36 to −10.62 ppm, widely shifted out of the diamagnetic region by proximity to the paramagnetic uranium centre. Taken together, the characterising data show that the carbodiimide has been coupled by the reducing U centre to form the rearranged cyclohexyl-substituted guanidinate ligand *via* loss of *cyclo*-hexylisonitrile. This presumably occurs similarly to the procedure reported for the reaction that converts $[\text{U}(\eta^5\text{-C}_5\text{Me}_5)_2(\eta^2\text{-bipy})]$ to $[\text{U}(\eta^5\text{-C}_5\text{Me}_5)_2\{\eta^2\text{-N}(\text{Cy})\text{C}(\text{=NCy})\text{N}(\text{Cy})\}]$ *via* extrusion of $[\text{CyNC}]$ from a coordinated DCC ligand then a $[2 + 2]$ addition to the transiently formed $\text{U}^{\text{V}}=\text{NCy}$ group.⁸¹ The fact that this forms from the U^{IV} product again supports the stabilising effect of this ligand on the higher formal oxidation states of uranium.

Reactions of **1^t** with CO and CO₂

In contrast to **1^m**, the bulkier **1^t** reduces both CO and CO₂; reaction with the former affording the known oxidation product **2^t**, and the latter affording a bridging carbonate **9^t**, although the yields are moderate and minimal respectively, Scheme 4.

Exposure of a degassed purple-brown solution of **1^t** in cyclopentane to a 1 atm pressure of CO₂ results in the immediate formation of a green solution. Colourless crystals can be obtained upon allowing the solution to stand at room temperature overnight. The single crystals collected were of sufficient quality to obtain connectivity information by single crystal XRD analysis to identify the product as $[(\mu\text{-CO}_3)\{\text{U}(\text{OBTrip}_2)_3\}_2]$ (**9^t**). The solid-state structure is reminiscent of other bimetallic carbonate-bridged uranium(IV) complexes,^{85,86} with a disordered bridging carbonate linking the uranium centres with distorted trigonal bipyramidal geometries. The boroxide

ligands are staggered along the U₁–C₁–U_{1A} axis to minimise the steric interactions of the ligands. The formation of carbonate bridged bimetallic uranium(IV) complexes has been postulated to proceed *via* a terminal uranium(V) oxo.^{87,88}

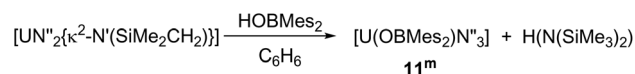
Comparatively, exposure of **1^t** solutions to a CO atmosphere only results in decomposition to the homoleptic U(IV) boroxide complex **2^t** and no CO containing products could be identified.

Formation of complexes $[\text{U}(\text{OBMe}_2)\text{N}''_3]$ **11^m** and $[\text{U}(\text{OBTrip}_2)\text{N}''_3]$ **11^t**

As mentioned previously, any traces of the metallacycle $[\text{UN}''_2\{\kappa^2\text{-N}'(\text{SiMe}_2\text{CH}_2)\}]$ in the synthesis of complexes **1^m** and **1^t** leads to the formation of complexes $[\text{UN}''_3(\text{OBMe}_2)]$ (**11^m**) and $[\text{UN}''_3(\text{OBTrip}_2)]$ (**11^t**). Moreover, an alternative synthesis for complex **11^m** is achieved by addition of just one equivalent of HOBMe₂ to an orange-brown solution of the uranium metallacycle $[\text{UN}''_2\{\kappa^2\text{-N}'(\text{SiMe}_2\text{CH}_2)\}]$, which provides a brown solution from which $[\text{UN}''_3(\text{OBMe}_2)]$ (**11^m**) can be isolated as brown solids (Scheme 5). Formation of complex **11^m** was confirmed by ¹H and ²⁹Si spectroscopies, elemental analysis and single crystal XRD analysis.

The ¹H NMR spectrum shows a paramagnetically shifted resonance at −6.56 ppm, which corresponds to the methyl groups in the tris(silylamide) ligand, and three smaller sets of resonances for the boroxide ligand. Moreover, the ²⁹Si NMR resonances shift from −74.46 and −81.84 ppm for the uranium metallacycle to −131.74 ppm for complex **11^m**.

Single crystal XRD analysis revealed that the solid-state structure of complexes **11^m** and **11^t** have a distorted tetrahedral coordination environment where the uranium centre is coordinated by one terminal boroxide ligand and three silylamide groups (Fig. 6). Crystals of complex **11^m** suitable for



Scheme 5 Synthesis of heteroleptic complex **11^m**.

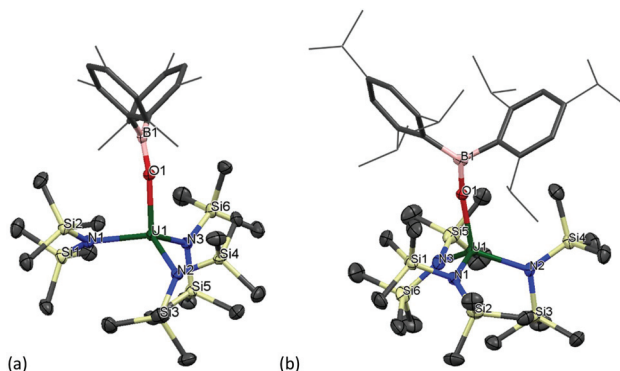
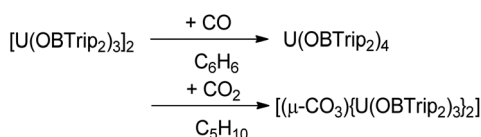


Fig. 6 Solid-state structure of (a) **11^m** and (b) **11^t**. Mesityl and triisopropylphenyl ligands are depicted as capped sticks and wireframe for clarity. Hydrogen atoms are omitted. The thermal ellipsoids are displayed at 50% probability.



Scheme 4 Reactions of complex **1^t** with CO and CO₂.



XRD analysis are obtained from a concentrated toluene solution. The average U–N_{avg} bond length is 2.268 Å, which is ~0.03 Å shorter than that of the previously reported complex [U(N(SiMe₃)₄)]⁸⁹ which is reasonable if more electron density is being distributed to the boroxide ligand and more electron density is being pulled from the tris(silylamide) ligands, thus decreasing the U–N bond length. The B–O_{avg} is 1.365(4) Å, which is longer than the B–O_{avg} bond length in complex **1^m** (mean 1.351 Å). Moreover, the N–U–N angles are 105.00(8)°, 113.72(8)° and 116.96(9)°, which are in the same range as for [U(N(SiMe₃)₄)]. The U–O distance is 2.1559(18) Å, which is slightly shorter than the U–O distance in the terminal boroxides for complex **1^m** (U–O_{avg} is 2.196 Å) but longer than the U–O_{avg} for **2^m** (2.144 Å). The U–O–B angle is 167.23(19)°, which is in the same range as U–O–B angles for the terminal boroxides in complex **1** (mean 171.6°). Colourless crystals of complex **11^t** suitable for X-ray diffraction analysis were obtained from a concentrated benzene solution upon standing at room temperature. The U–O distance amounts to 2.167(3) Å, which is longer than that of OB(Mes)₂-coordinating analogue **11^m**. The mean U–N bond length is 2.272(3) Å, which is longer than complex **11^m** and the previously reported [U(N(SiMe₃)₄)]⁸⁹. Bond length of B–O (1.365(5) Å) is essentially the same as that of U(IV) complex **2^t** (1.364(7) Å). Bond angle of U–O–B is 169.8(3)°, which is in the same range as the U–O–B angle in complex **11^m**.

Conclusions

Diaryl boroxide ligands are capable ligands for both U^{III} and U^{IV} cations but the U^{III} complexes are particularly air-sensitive and decompose readily to different U^{IV} complexes depending on the conditions. We suggest that rather than working in tandem as a second Lewis acidic reaction centre, the role of the boron is simply to remove electron density from the reducing U^{III} centre in **1** to such an extent that it does not efficiently reduce an incoming substrate, and neither the U or B centres in these complexes can bind a phosphine. No reactions were observed for **1^m** with carbon monoxide or carbon dioxide at ambient pressure and temperature, underlining these observations. Reactions at elevated temperatures simply result in the decomposition of **1^m**. Meanwhile, with larger boroxide aryl substituents, complex **1^t** exhibits greater thermal stability compared to less encumbered **1^m** and greater reactivity with the less reducible, smaller carbon oxides, but lower reactivity with the less reactive, but slightly larger small molecules tested so far. In general, the reactions of **1^m** with small molecules and other substrates give low yields due to the extensive formation of **3^m** from traces of oxygen, or perhaps also reactions with vacuum grease and/or glassware. Qualitatively, the sum of these reactions suggest that the OBAR₂ ligands make the UX₃ centre less strongly reducing than the analogous aryloxide ligands ODtbp, which is in line with expectation, but we caution that due to the additional modes of reactivity available to the boron centre, a direct and more precise estimate of the

relative oxidation potentials that the ligands impose on the U(III) centre in each complex is more difficult to make. For example, the aryloxide complexes U(OAr)₃ reductively activate N₂ but this has not been observed in the boroxide system, although the increased capacity for both OBAR₂ ligands to bridge to a second U centre hampers the comparison. Work on finding a more quantitative measure for this is in progress. The isolation of small quantity of a μ -carbonate product [(μ -CO₃){U(OBTrip₂)₃]₂] suggests there are opportunities however, and work is in progress to identify new substrates that could be usefully activated by the combination of the two potentially Lewis acidic centres in this system.

Experimental section

[U(OBMes₂)₃]₂ (**1^m**)

A Schlenk flask was charged with U[N(SiMe₃)₂]₃ (1.08 g, 1.5 mmol) and HOBMes₂ (1.20 g, 4.5 mmol, 3 eq.) and a stir bar. The reaction mixture was dissolved in toluene (40 mL) to yield a purple solution, which was allowed to stir at room temperature for 4 hours. Volatiles were removed by evaporation to dryness. The residue was washed with cold hexane (–78 °C, 4 mL) and recrystallized from hexane providing dark purple block crystals suitable for single-crystal X-ray crystallography. Yield: 634 mg, 41%.

¹H NMR (500 MHz, C₆D₆): δ_{H} 8.29 (Ar-H terminal Mes, 16H); 7.20 (*o*-CH₃ terminal Mes, 48H); 2.53 (*p*-CH₃ terminal Mes, 24H); –3.71 (*o*-CH₃ bridging Mes, 12H); –9.20 (Ar-H and *p*-CH₃ bridging Mes, 20H); –15.13 (*o*-CH₃ bridging Mes, 12H).

¹¹B NMR (160.46 MHz, C₆D₆): δ_{B} 73.65 (OBMes). The latter is a tentative assignment. The sample analysed was dissolved single crystals, but this resonance is close to that of hydrolysed ligand so it may be the other resonance is not observed in the window.

MS (APPI): m/z = 1033 ([M]⁺ [U(OBMes₂)₃]) C₅₄H₆₆B₃O₃U⁺ [M]⁺ requires 1033.5799, found 1033.5675 (+12.4 ppm).

FTIR (cm^{–1}): 3612, 2728, 1721, 1608, 1552, 1280, 1228, 1176, 1151, 1081, 1029, 960, 928, 847, 745, 672.

Anal. calc for C₁₀₄H₁₃₂B₆O₆U₂: C, 62.75 H, 6.44 N, 0.00%. Found: C, 62.63 H, 6.47 N, 0.00%.

[U(OBMes₂)₄] (**2^m**)

A toluene solution (1.5 mL) of NaOBMes₂ (115 mg, 0.4 mmol, 4 eq.) was added dropwise over 10 min to a solution of UI₄(dioxane)₂ (92 mg, 0.1 mmol) in toluene (1.5 mL) in a 7 cm³ vial with stirring for 18 hours. The reaction was centrifuged and filtered. Colourless plates suitable for single-crystal X-ray crystallography were grown from slow diffusion of hexanes into the reaction mixture. Yield: 16.7 mg, 12%.

¹H NMR (500 MHz, C₆D₆): δ_{H} 6.71 (Ar-H Mes, 16H); 2.15 (*o*-CH₃ Mes, 48H); 1.99 (*p*-CH₃ Mes, 24H). ¹¹B NMR (160.46 MHz, C₆D₆): not observed in the window +130––130 ppm.



MS (APPI): m/z = 1299 ($[M]^{++}$ $[U(OBMeS_2)_4]$). $C_{72}H_{88}B_4O_4U$ $[M]^{++}$ requires 1298.7563, found 1298.7595 (−3.2 ppm).

$[U(OBTrip_2)_3]_2$ (1^t) $[U(OBTrip_2)_4]$ (2^t)

HOBTrip₂ (104 mg, 0.24 mmol) and UN³ (72 mg, 0.1 mmol) were dissolved in benzene (0.8 mL) in a 7 mL vial. The resulting dark red solution was stored at room temperature for 60 h during which time $U(OBTrip_2)_4$ 2^t crystallised as dark red crystals which can be isolated and washed with benzene (0.3 mL × 3). A second crop of 2^t was isolated from the mother liquor to give a combined yield of 25% of 2^t (30 mg, 0.02 mmol). Complex 1^t (39 mg, 0.013 mmol, 32% yield) was isolated after removal of volatiles from the mother liquor.

Characterization of complex 1^t :

1H NMR (500 MHz, C_6D_6): δ 8.10 (br-s, 12H, Ar-H), 2.90 (br-s, 18H, CH), 1.45 (br-s, 108H, Me). ^{11}B NMR (160.46 MHz, C_6D_6): 110.19 (O-B-Ar). Repeated submissions of samples for elemental analysis did not yield reasonable data.

Characterization of complex 2^t :

1H NMR (500 MHz, C_6D_6): δ 44.14 (s, 3H, Ar-H), 17.87 (s, 3H, Ar-H), 10.66 (s, 4H, Ar-H), 8.86 (s, 6 Ar-H + 4 CH), 7.41 (s, 8H, CH), 6.74 (s, 4H, CH), 5.46 (s, 4H, CH), 4.78 (s, 4H, CH), 4.13 (s, 12H, CH₃), 1.49 (m, 24H, CH₃), 1.24 (s, 24H, CH₃), 0.52 (d, J = 6 Hz, 12H, CH₃), 0.25 (d, J = 6 Hz, 12H, CH₃), 0.02 (d, J = 6 Hz, 9H, CH₃), −1.34 (s, 3H, CH₃), −3.43 (s, 12H, CH₃), −4.29 (s, 12H, CH₃), −13.75 (s, 12H, CH₃), −16.19 (s, 9H, CH₃), −18.40 (s, 3H, CH₃). ^{11}B NMR (160.46 MHz, C_6D_6): no obvious resonance in the range of 230–40 ppm. Repeated submissions of samples for elemental analysis did not yield reasonable data.

$[U(OBTrip_2)_3(OEt_2)]$ (1^t -OEt₂)

A solution of HOBTrip₂ (326 mg, 0.75 mmol) in Et₂O (1 mL) was added dropwise to solution of UN³ (216 mg, 0.3 mmol) in Et₂O (1 mL) in a 7 mL vial. The resulting dark red solution was stirred at room temperature for 0.5 h, which was left at −30 °C for a further *ca.* 60 h. A brown precipitate was separated and washed with Et₂O (0.5 mL × 3) as complex 1^t (see below). The dark red mother liquid was concentrated, and left for standing, which gave complex 1^t -OEt₂ (46 mg, 0.029 mmol) as dark-red crystals. The isolated yield is low due to the very high solubility of the product.

1H NMR (500 MHz, C_6D_6): δ 8.94 (br, 12H, Ar-H), 3.56 (br-s, 18H, CH), 1.73 (br-s, 108H, Me). ^{11}B NMR (160.46 MHz, C_6D_6): not observed in the window +130–−130 ppm. No reasonable elemental analysis data were obtained despite repeated trials.

$[OU(OBMeS_2)_3]_n$ (3^m)

Method A. This product is observed when the complex $[U(OBMeS_2)_2(\mu-OBMeS_2)]_2$ is stored as a solution or when solutions are heated, due to reaction with traces of oxygen.

Method B. A purple solution of $[U(OBMeS_2)_2(\mu-OBMeS_2)]_2$ (25 mg, 0.012 mmol) in C_6D_6 (0.4 mL) was prepared *in situ* in a Young's NMR tube. The solution was then degassed by three freeze pump thaw cycles and exposed to 1 bar pressure of N₂O. The mixture was agitated to give an immediate colour change

to light brown. Formation of complex $[OU(OBMeS_2)_3]_n$ was confirmed by NMR spectroscopy and elemental analysis.

1H NMR (500 MHz, C_6D_6): δ_H 5.90 (Ar-H Mes); 1.89 (*p*-CH₃ Mes); −0.52 (*o*-CH₃ Mes).

^{11}B NMR (160.46 MHz, C_6D_6): δ_B 89.29 (OBMeS).

^{13}C NMR (126 MHz, C_6D_6): δ_C 184.55 (free CO); 166.10 (*o*-C Mes); 137.10 (*p*-C Mes); 133.59 (C-B Mes); 126.94 (Ar-CH Mes); 21.29 (*p*-CH₃ Mes); 5.05 (*o*-CH₃ Mes).

Elemental analysis: C 61.80%, H 6.34% calculated. C 61.75%, H 6.37% found.

FTIR (cm^{−1}): 3617, 2728, 1721, 1608, 1552, 1280, 1228, 1175, 1152, 1081, 1029, 960, 928, 840, 745, 670.

$[U(OBMeS_2)_4(OPy)_2]_n$ (4^m)

A Young's NMR tube was charged with $[U(OBMeS_2)_2(\mu-OBMeS_2)]_2$ (25 mg, 0.012 mmol), C₅H₅NO (5 mg, 0.048 mmol, 4 eq.) and C_6D_6 (0.4 mL). The reaction mixture turned dark brown immediately and was allowed to react for 72 hours, during which time dark orange crystals of $[U(OBMeS_2)_4(Py-O)_2]$ deposited on the reaction vessel walls. Yield: 19 mg, 65%.

1H NMR (500 MHz, C_6D_6): δ_H 6.76 (Ar-H Mes, 12H); 4.42 (Py-O H, 6H); 2.31 (*o*-CH₃ Mes, 36H); 2.16 (*p*-CH₃ Mes, 18H) 1.96 (Py-O H, 4H).

^{11}B NMR (160.46 MHz, C_6D_6): not observed in the window +130–−130 ppm.

MS (APPI): m/z = 1223 ($[M]^{++}$ $[U(OBMeS_2)_3(Py-O)_2]$). $C_{64}H_{76}B_3N_2O_5U^{++}$ $[M]^{++}$ requires 1223.6541, found 1223.6552 (−1.1 ppm).

Dark orange crystals suitable for SCXRD analysis were grown from concentrated benzene solutions, however, the data are only of sufficient quality to discuss connectivity and not detailed metrics.

$[U(OBMeS_2)_3(OPPh_3)_2]$ (5^m and 6^m)

$[U(OBMeS_2)_2(\mu-OBMeS_2)]_2$ (25 mg, 0.012 mmol) and OPPh₃ (7 mg, 0.024 mmol, 2 eq.) were placed in a Young's NMR tube. The tube's contents were dissolved in C_6D_6 (0.4 mL). 1H NMR confirmed the formation of $[U(OBMeS_2)_3(OPPh_3)_2]$ (5^m). Co-crystals of 5^m and the side product $[U(OBMeS_2)_3(B_2O_3MeS_2)]$ (6^m) suitable for X-ray diffraction studies were grown by slow diffusion of hexane into a concentrated benzene solution of $[U(OBMeS_2)_3(OPPh_3)_2]$. Yield: 23 mg, 60%.

1H NMR (500 MHz, C_6D_6): δ_H 6.90 (Ar-H Mes, 12H); 6.46 (*p*-Ph H, 6H); 5.92 (*o*-Ph H, 12H); 3.64 (*o*-CH₃ Mes, 36H); 2.85 (*m*-Ph H, 12H); 1.87 (*p*-CH₃ Mes, 18H).

^{11}B NMR (160.46 MHz, C_6D_6) (5^m): δ_B not observed in the window +130–−130 ppm.

^{31}P NMR (202 MHz, C_6D_6): δ_P −17.83(OPPh₃).

MS (APPI): m/z = 1589 ($[M]^{++}$ $[U(OBMeS_2)_3(OPPh_3)_2]$). $C_{90}H_{96}B_3O_5P_2U^{++}$ $[M]^{++}$ requires 1589.7520, found 1589.7790 (−27.0 ppm).

$[U(OBMeS_2)_3]_2(\mu-\eta^2:\eta^2-S_2)_2$ (7^m)

To a magnetically stirred, dark purple solution of $[U(OBMeS_2)_2(\mu-OBMeS_2)]_2$ (165 mg, 0.08 mmol) in 2 mL of toluene in a 7 cm³ vial was added elemental sulfur (10 mg,



0.04 mmol, 0.5 eq.) with stirring and the reaction was allowed to stir for 2 hours. Dark red crystals of $[\{(\text{OBMe}_2)_3\text{U}\}_2(\mu-\eta^2:\eta^2-\text{S}_2)_2]$ suitable for X-ray diffraction can be obtained from storage of a concentrated benzene solution at room temperature. Yield: 32 mg, 18%.

^1H NMR (500 MHz, C_6D_6): δ_{H} 6.88 (Ar-H Mes, 24H); 2.38 (*p*-CH₃ Mes, 36H); 2.23 (*o*-CH₃ Mes, 72H).

^{11}B NMR (160.46 MHz, C_6D_6): not observed in the window +130–130 ppm. MS (APPI): m/z = 1097 ($[\text{M}]^{+}$). $[\text{U}(\text{OBMe}_2)_3(\mu-\text{S}_2)] \text{C}_{54}\text{H}_{66}\text{B}_3\text{O}_3\text{S}_2\text{U}^{+} [\text{M}]^{+}$ requires 1097.5240, found 1097.5259 (−1.9 ppm).

$[\text{U}(\eta^2\text{-DCC})(\text{OBMe}_2)_3] (\mathbf{8}^{\text{m}})$

A solution of *N,N'*-dicyclohexylcarbodiimide [$\text{DCC} = \eta^2\text{-N}(\text{Cy})\text{C}(\text{=N}(\text{Cy}))\text{N}(\text{Cy})$; Cy = C_6H_{11}] (10 mg, 0.048 mmol, 2 eq.) in benzene (1 mL) was added dropwise to a stirring benzene solution of $[\text{U}(\text{OBMe}_2)_2(\mu\text{-OBMe}_2)]_2$ (50 mg, 0.024 mmol) in a 7 cm³ vial. The solution turned green immediately and was allowed to stir for 18 hours. The solution was then filtered and evaporated to dryness. ^1H NMR spectroscopy confirmed the formation of the title compound. Yield: 22 mg, 69%.

^1H NMR (500 MHz, C_6D_6): δ_{H} 58.36 (Cy H, 1H); 43.14 (Cy H, 1H); 28.62 (Cy H, 3H); 13.73 (Cy H, 2H); 12.48 (Cy H, 1H); 11.15 (Cy H, 2H); 9.66 (Cy H, 1H); 9.06 (Cy H, 3H); 7.64 (Cy H, 2H); 7.32 (Cy H, 1H); 6.75 (Ar-H Mes, 12H); 6.48 (Cy H, 2H); 3.89 (Cy H, 1H); 3.08 (*p*-CH₃ Mes, 18H); 2.92 (Cy H, 4H); 2.55 (Cy H, 1H); 2.16 (Cy H, 1H); 1.76 (Cy H, 1H); −4.75 (Cy H, 1H); −6.63 (Cy H, 1H); −10.39 (*o*-CH₃ Mes, 36H); −10.62 (Cy H, 4H).

MS (APPI): m/z = 1338 ($[\text{M} + \text{H}]$) $(\text{OBMe}_2)_3\text{U}[\eta^2\text{-N}(\text{C}_6\text{H}_{11})\text{C}(\text{=NC}_6\text{H}_{11})\text{N}(\text{C}_6\text{H}_{11})] \text{C}_{73}\text{H}_{100}\text{B}_3\text{N}_3\text{O}_3\text{U}^{+} [\text{M} + \text{H}]^{+}$ requires 1337.8552, found 1337.8577 (−2.5 ppm).

$[\text{U}(\text{OBTrip}_2)_3]_2(\mu\text{-CO}_3) (\mathbf{9}^{\text{f}})$

A purple-brown solution of $[\text{U}(\text{OBTrip}_2)_3]_2$ (75 mg, 0.048 mmol, 1 eq.) in C_5H_{10} prepared *in situ* in a Young's NMR tube was degassed by three freeze pump thaw cycles and placed under a 1 bar pressure of CO₂ at room temperature. The mixture was agitated to give an immediate colour change to light green-brown. The reaction mixture was allowed to stand overnight at room temperature. Crystalline $[\text{U}(\text{OBTrip}_2)_3]_2(\mu\text{-CO}_3)$ was obtained by slow diffusion of hexanes into the reaction mixture. Yield: 4 mg, 2%. Due to the extremely low yield of isolated material, no further characterisation data were obtainable.

$[\text{U}(\text{OBMe}_2)_3\text{N}''_3] (\mathbf{11}^{\text{m}})$

Method A: UIN''_3 (225 mg) and KOBMe_2 (100 mg) were placed into a Schlenk flask with a stirrer bar. Toluene (15 mL) was introduced into the flask and the contents were stirred overnight. The solution was then cannula filtered and the solvent removed. The resulting residue was washed with cold hexanes (2 × 5 mL) to yield the title compound $\mathbf{11}^{\text{m}}$ (205 mg, 78%).

Method B: $(\text{N}'')_2\text{U}\{\kappa^2\text{-N}(\text{C}_6\text{H}_5)_2\text{N}(\text{SiMe}_3)_2\text{SiMe}_2\text{CH}_2\}$ (10 mg) and dimesitylborinic acid (3.5 mg) were placed in an NMR tube and dissolved in C_6D_6 (0.5 mL). ^1H NMR showed that quantitative formation of $[\text{U}(\text{OBMe}_2)_3\text{N}''_3]$ had occurred. Crystals suit-

able for X-ray diffraction studies were grown by slow evaporation of a toluene solution of $[\text{U}(\text{OBMe}_2)_3\text{N}''_3]$. Local code P18053_tri.

^1H NMR (C_6D_6): δ_{H} 7.52 (s, 4 H), 4.89 (s, 12 H), 2.48 (s, 6 H), −6.68 (s, 54 H); anal. Calc for $\text{C}_{36}\text{H}_{76}\text{B}_1\text{N}_3\text{O}_1\text{Si}_6\text{U}_1$: C, 52.58 H, 8.01 N, 3.41%. Found: C, 52.39 H, 7.34 N, 3.53%; EI-MS(m/z): 823.2 $[\text{M} - \text{N}''_3]^+$.

$[\text{U}(\text{OBTrip}_2)_3\text{N}''_3] (\mathbf{11}^{\text{t}})$

Complex $[\text{U}(\text{OBTrip}_2)_3\text{N}''_3]$ $\mathbf{11}^{\text{t}}$ was also isolated as a side product in 5% yield from the synthesis of $\mathbf{1}^{\text{t}}\text{-OEt}_2$ as a brown precipitate, and washed with Et₂O (0.5 mL × 3) in low yield (46 mg, 0.04 mmol, 5% yield based on HOBTrip_2). Colourless crystals of $\mathbf{11}^{\text{t}}$ suitable for X-ray diffraction analysis were obtained from concentrated benzene solution upon standing at room temperature. Local code P18083.

^1H NMR (500 MHz, C_6D_6): δ 41.93 (s, 1H, Ar-H), 16.81 (s, 1H, Ar-H), 11.70 (s, 2H, Ar-H), 10.47 (s, 1H, Ar-H), 9.29 (s, 2H, CH), 8.01 (s, 1H, CH), 7.48 (s, 2H, CH), 3.25 (s, 12H, CH₃), 1.98 (s, 3H, CH₃), 1.21 (s, 3H, CH₃), 1.10 (s, 3H, CH₃), 0.52 (s, 3H, CH₃), 1.48 (s, 9H, CH₃), −4.63 (s, 3H, CH₃), −7.36 (s, 12H, CH₃), −10.03 (s, 24H, CH₃), −14.56 (s, 18H, CH₃). There is restricted rotation of the CH₃ groups at room temperature on the ^1H NMR timescale. ^{11}B NMR (160.46 MHz, C_6D_6): 109.74 (O–B–Ar).

Conflicts of interest

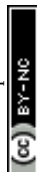
There are no conflicts to declare.

Acknowledgements

We thank the University of Edinburgh and the EPSRC for funding through EP/H004823/1 and EP/M010554/1, EP/J018139/1 and the UK Catalysis Hub EP/K014714/1. This project has also received funding from the European Research Council (ERC) under the European Union's Horizon 2020 research and innovation programme (grant agreement No 740311). The support provided by China Scholarship Council (CSC) during a visit of D. Y. to the University of Edinburgh is acknowledged. We would also like to thank Dr Colin Logan Mackay of the University of Edinburgh for mass spectrometry and the University of Edinburgh SIRCAMS facility for use of the FTICRMS.

References

- 1 P. L. Arnold, Z. R. Turner, R. M. Bellabarba and R. P. Tooze, *Chem. Sci.*, 2011, **2**, 77–79.
- 2 S. M. Mansell, N. Kaltsoyannis and P. L. Arnold, *J. Am. Chem. Soc.*, 2011, **133**, 9036–9051.
- 3 S. M. Mansell, J. H. Farnaby, A. I. Germeroth and P. L. Arnold, *Organometallics*, 2013, **32**, 4214–4222.



- 4 M. Falcone, L. Chatelain, R. Scopelliti, I. Živković and M. Mazzanti, *Nature*, 2017, **547**, 332.
- 5 V. Mougél, C. Camp, J. Pécaut, C. Copéret, L. Maron, C. E. Kefalidis and M. Mazzanti, *Angew. Chem., Int. Ed.*, 2012, **51**, 12280–12284.
- 6 P. L. Arnold, S. M. Mansell, L. Maron and D. McKay, *Nat. Chem.*, 2012, **4**, 668–674.
- 7 S. C. Cole, M. P. Coles and P. B. Hitchcock, *J. Chem. Soc., Dalton Trans.*, 2002, 4168–4174.
- 8 M. P. Coles, *Coord. Chem. Rev.*, 2016, **323**, 52–59.
- 9 O. J. Cooper, J. McMaster, W. Lewis, A. J. Blake and S. T. Liddle, *Dalton Trans.*, 2010, **39**, 5074–5076.
- 10 D. M. King, W. Lewis and S. T. Liddle, *Inorg. Chim. Acta*, 2012, **380**, 167–173.
- 11 I. Korobkov, S. Gorelsky and S. Gambarotta, *J. Am. Chem. Soc.*, 2009, **131**, 10406–10420.
- 12 C. R. Groom, I. J. Bruno, M. P. Lightfoot and S. C. Ward, *Acta Crystallogr., Sect. B: Struct. Sci., Cryst. Eng. Mater.*, 2016, **72**, 171–179.
- 13 A.-C. Schmidt, F. W. Heinemann, W. W. Lukens and K. Meyer, *J. Am. Chem. Soc.*, 2014, **136**, 11980–11993.
- 14 M. J. Monreal, R. K. Thomson, T. Cantat, N. E. Travia, B. L. Scott and J. L. Kiplinger, *Organometallics*, 2011, **30**, 2031–2038.
- 15 J. A. L. Wells, M. L. Seymour, M. Suvova and P. L. Arnold, *Dalton Trans.*, 2016, **45**, 16026–16032.
- 16 J. A. L. Wells, PhD thesis, University of Edinburgh, 2018.
- 17 A. Zalkin, J. G. Brennan and R. A. Andersen, *Acta Crystallogr., Sect. C: Cryst. Struct. Commun.*, 1988, **44**, 1553–1554.
- 18 C. J. Burns, W. H. Smith, J. C. Huffman and A. P. Sattelberger, *J. Am. Chem. Soc.*, 1990, **112**, 3237–3239.
- 19 C. Camp, J. Pécaut and M. Mazzanti, *J. Am. Chem. Soc.*, 2013, **135**, 12101–12111.
- 20 H. J. Wasserman, D. C. Moody and R. R. Ryan, *J. Chem. Soc., Chem. Commun.*, 1984, 532–533.
- 21 D. S. J. Arney and C. J. Burns, *J. Am. Chem. Soc.*, 1995, **117**, 9448–9460.
- 22 D. S. J. Arney and C. J. Burns, *J. Am. Chem. Soc.*, 1993, **115**, 9840–9841.
- 23 O. Cooper, C. Camp, J. Pécaut, C. E. Kefalidis, L. Maron, S. Gambarelli and M. Mazzanti, *J. Am. Chem. Soc.*, 2014, **136**, 6716–6723.
- 24 S. J. Kraft, J. Walensky, P. E. Fanwick, M. B. Hall and S. C. Bart, *Inorg. Chem.*, 2010, **49**, 7620–7622.
- 25 L. R. Avens, D. M. Barnhart, C. J. Burns, S. D. McKee and W. H. Smith, *Inorg. Chem.*, 1994, **33**, 4245–4254.
- 26 A. J. Lewis, U. J. Williams, J. M. Kikkawa, P. J. Carroll and E. J. Schelter, *Inorg. Chem.*, 2012, **51**, 37–39.
- 27 N. L. Bell, P. L. Arnold and J. B. Love, *Dalton Trans.*, 2016, **45**, 15902–15909.
- 28 J. A. Pool, B. L. Scott and J. L. Kiplinger, *J. Am. Chem. Soc.*, 2005, **127**, 1338–1339.
- 29 N. A. Siladke, J. LeDuc, J. W. Ziller and W. J. Evans, *Chem. – Eur. J.*, 2012, **18**, 14820–14827.
- 30 J.-C. Berthet, M. Nierlich and M. Ephritikhine, *Dalton Trans.*, 2004, 2814–2821.
- 31 S. Kannan, M. A. Moody, C. L. Barnes and P. B. Duval, *Inorg. Chem.*, 2006, **45**, 9206–9212.
- 32 J. J. Kiernicki, J. S. Harwood, P. E. Fanwick and S. C. Bart, *Dalton Trans.*, 2016, **45**, 3111–3119.
- 33 T. W. Hayton, J. M. Boncella, B. L. Scott, E. R. Batista and P. J. Hay, *J. Am. Chem. Soc.*, 2006, **128**, 10549–10559.
- 34 R. E. Jilek, L. P. Spencer, R. A. Lewis, B. L. Scott, T. W. Hayton and J. M. Boncella, *J. Am. Chem. Soc.*, 2012, **134**, 9876–9878.
- 35 R. E. Jilek, N. C. Tomson, B. L. Scott and J. M. Boncella, *Inorg. Chim. Acta*, 2014, **422**, 78–85.
- 36 J.-C. Berthet, M. Nierlich and M. Ephritikhine, *Polyhedron*, 2003, **22**, 3475–3482.
- 37 N. A. Ayoub, A. R. Browne, B. L. Anderson and T. G. Gray, *Dalton Trans.*, 2016, **45**, 3820–3830.
- 38 Z. Yang, X. Ma, R. B. Oswald, H. W. Roesky and M. Noltemeyer, *J. Am. Chem. Soc.*, 2006, **128**, 12406–12407.
- 39 X. Ma, Z. Yang, X. Wang, H. W. Roesky, F. Wu and H. Zhu, *Inorg. Chem.*, 2011, **50**, 2010–2014.
- 40 Z. Yang, P. Hao, Z. Liu, X. Ma, H. W. Roesky and J. Li, *J. Organomet. Chem.*, 2014, **751**, 788–791.
- 41 M. Kořenková, B. Mairychová, R. Jambor, Z. Růžicková and L. Dostál, *Inorg. Chem. Commun.*, 2014, **47**, 128–130.
- 42 M. Kořenková, M. Erben, R. Jambor, A. Růžicka and L. Dostál, *J. Organomet. Chem.*, 2014, **772–773**, 287–291.
- 43 B. Mairychová, T. Svoboda, P. Štěpnička, A. Růžicka, R. W. A. Havenith, M. Alonso, F. D. Proft, R. Jambor and L. Dostál, *Inorg. Chem.*, 2013, **52**, 1424–1431.
- 44 B. Mairychová, P. Štěpnička, A. Růžicka, L. Dostál and R. Jambor, *Organometallics*, 2014, **33**, 3021–3029.
- 45 B. Mairychová, I. V. Kityk, A. Maciag, F. Bureš, M. Klikar, A. Růžicka, L. Dostál and R. Jambor, *Inorg. Chem.*, 2016, **55**, 1587–1594.
- 46 L. Dostál, R. Jambor, A. Růžicka, R. Jirásko, A. Lyčka, J. Beckmann and S. Ketkov, *Inorg. Chem.*, 2015, **54**, 6010–6019.
- 47 M. Kořenková, B. Mairychová, A. Růžicka, R. Jambor and L. Dostál, *Dalton Trans.*, 2014, **43**, 7096–7108.
- 48 U. Bossek, H. Hummel, T. Weyhermüller, K. Wieghardt, S. Russell, L. van der Wolf and U. Kolb, *Angew. Chem., Int. Ed.*, 1996, **35**, 1552–1554.
- 49 A. R. Browne, N. Deligonul, B. L. Anderson, M. Zeller, A. D. Hunter and T. G. Gray, *Chem. Commun.*, 2015, **51**, 15800–15803.
- 50 I. Pantcheva and K. Osakada, *Organometallics*, 2006, **25**, 1735–1741.
- 51 H. Yin, A. J. Lewis, U. J. Williams, P. J. Carroll and E. J. Schelter, *Chem. Sci.*, 2013, **4**, 798–805.
- 52 E. Barnea, T. Andrea, M. Kapon and M. S. Eisen, *J. Am. Chem. Soc.*, 2004, **126**, 5066–5067.
- 53 R. K. Das, E. Barnea, T. Andrea, M. Kapon, N. Fridman, M. Botoshansky and M. S. Eisen, *Organometallics*, 2015, **34**, 742–752.
- 54 K. I. M. Ingram, N. Kaltsoyannis, A. J. Gaunt and M. P. Neu, *J. Alloys Compd.*, 2007, **444–445**, 369–375.
- 55 H. H. Dam, D. N. Reinhoudt and W. Verboom, *Chem. Soc. Rev.*, 2007, **36**, 367–377.



- 56 S. R. Daly, J. M. Keith, E. R. Batista, K. S. Boland, D. L. Clark, S. A. Kozimor and R. L. Martin, *J. Am. Chem. Soc.*, 2012, **134**, 14408–14422.
- 57 D. E. Smiles, G. Wu, P. Hrobárik and T. W. Hayton, *J. Am. Chem. Soc.*, 2016, **138**, 814–825.
- 58 J. K. Pagano, D. S. J. Arney, B. L. Scott, D. E. Morris, J. L. Kiplinger and C. J. Burns, *Dalton Trans.*, 2019, **48**, 50–57.
- 59 S. M. Franke, F. W. Heinemann and K. Meyer, *Chem. Sci.*, 2014, **5**, 942–950.
- 60 B. M. Gardner, D. M. King, F. Tuna, A. J. Wooles, N. F. Chilton and S. T. Liddle, *Chem. Sci.*, 2017, **8**, 6207–6217.
- 61 D. E. Smiles, G. Wu and T. W. Hayton, *New J. Chem.*, 2015, **39**, 7563–7566.
- 62 D. L. Perry, A. Zalkin, H. Ruben and D. H. Templeton, *Inorg. Chem.*, 1982, **21**, 237–240.
- 63 D. J. Grant, Z. Weng, L. J. Jouffret, P. C. Burns and L. Gagliardi, *Inorg. Chem.*, 2012, **51**, 7801–7809.
- 64 J. L. Brown, G. Wu and T. W. Hayton, *Organometallics*, 2013, **32**, 1193–1198.
- 65 E. M. Matson, M. D. Goshert, J. J. Kiernicki, B. S. Newell, P. E. Fanwick, M. P. Shores, J. R. Walensky and S. C. Bart, *Chem. – Eur. J.*, 2013, **19**, 16176–16180.
- 66 A. C. Sutorik and M. G. Kanatzidis, *Polyhedron*, 1997, **16**, 3921–3927.
- 67 J.-E. Kwak, D. L. Gray, H. Yun and J. A. Ibers, *Acta Crystallogr., Sect. E: Struct. Rep. Online*, 2006, **62**, i86–i87.
- 68 C. Camp, M. A. Antunes, G. García, I. Ciofini, I. C. Santos, J. Pécaut, M. Almeida, J. Marçalo and M. Mazzanti, *Chem. Sci.*, 2014, **5**, 841–846.
- 69 A. Terzis and R. Rivest, *Inorg. Chem.*, 1973, **12**, 2132–2136.
- 70 R. C. Elder and M. Trkula, *Inorg. Chem.*, 1977, **16**, 1048–1051.
- 71 J. T. York, E. C. Brown and W. B. Tolman, *Angew. Chem., Int. Ed.*, 2005, **44**, 7745–7748.
- 72 S. Yao, C. Milsmann, E. Bill, K. Wieghardt and M. Driess, *J. Am. Chem. Soc.*, 2008, **130**, 13536–13537.
- 73 W. J. Evans, J. R. Walensky, J. W. Ziller and A. L. Rheingold, *Organometallics*, 2009, **28**, 3350–3357.
- 74 W. J. Evans, M. K. Takase, J. W. Ziller and A. L. Rheingold, *Organometallics*, 2009, **28**, 5802–5808.
- 75 M. K. Takase, N. A. Siladke, J. W. Ziller and W. J. Evans, *Organometallics*, 2011, **30**, 458–465.
- 76 O. J. Cooper, D. P. Mills, W. Lewis, A. J. Blake and S. T. Liddle, *Dalton Trans.*, 2014, **43**, 14275–14283.
- 77 M. A. Boreen, B. F. Parker, T. D. Lohrey and J. Arnold, *J. Am. Chem. Soc.*, 2016, **138**, 15865–15868.
- 78 W. J. Evans, J. R. Walensky and J. W. Ziller, *Organometallics*, 2010, **29**, 101–107.
- 79 W. J. Evans, J. R. Walensky and J. W. Ziller, *Inorg. Chem.*, 2010, **49**, 1743–1749.
- 80 N. A. Siladke, J. W. Ziller and W. J. Evans, *J. Am. Chem. Soc.*, 2011, **133**, 3507–3516.
- 81 L. Zhang, C. Zhang, G. Hou, G. Zi and M. D. Walter, *Organometallics*, 2017, **36**, 1179–1187.
- 82 L. Zhang, G. Hou, G. Zi, W. Ding and M. D. Walter, *J. Am. Chem. Soc.*, 2016, **138**, 5130–5142.
- 83 L. Zhang, B. Fang, G. Hou, G. Zi, W. Ding and M. D. Walter, *Organometallics*, 2017, **36**, 898–910.
- 84 G. Zi, *Chem. Commun.*, 2018, **54**, 7412–7430.
- 85 O. T. Summerscales, A. S. P. Frey, F. G. N. Cloke and P. B. Hitchcock, *Chem. Commun.*, 2008, 198–200.
- 86 C. J. Inman, A. S. P. Frey, A. F. R. Kilpatrick, F. G. N. Cloke and S. M. Roe, *Organometallics*, 2017, **36**, 4539–4545.
- 87 N. Tsoureas, L. Castro, A. F. R. Kilpatrick, F. G. N. Cloke and L. Maron, *Chem. Sci.*, 2014, **5**, 3777–3788.
- 88 L. Castro, O. P. Lam, S. C. Bart, K. Meyer and L. Maron, *Organometallics*, 2010, **29**, 5504–5510.
- 89 A. J. Lewis, U. J. Williams, P. J. Carroll and E. J. Schelter, *Inorg. Chem.*, 2013, **52**, 7326–7328.

