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# Temperature induced single-crystal to single-crystal transformation of uranium azide complexes†

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The monomeric and dimeric uranium azide complexes  $\{[(CH_3)_2NCH_2-CH_2NPPr_2]_2U(N_3)_2\}$  (**2**) and  $\{[(CH_3)_2NCH_2CH_2NPPr_2]_2U(N_3)_2\}$  (**3**) were synthesized by treating complex **1** with  $NaN_3$  at 60 and  $-20^\circ C$ , respectively. A temperature-induced single-crystal to single-crystal transformation of **3** to **2** was observed. The reduction of either **2** or **3** with  $KC_8$  yields a uranium nitride complex  $\{[(CH_3)_2NCH_2CH_2NPPr_2]_4U_2(\mu-N)_2\}$  (**4**).

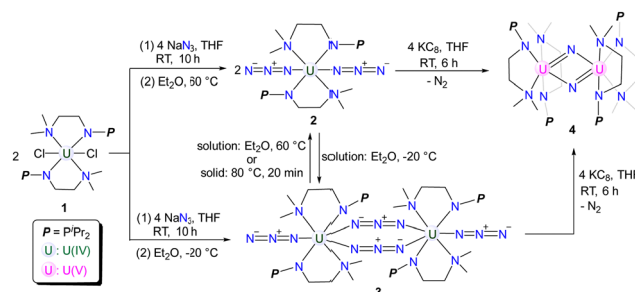
Single-crystal to single-crystal (SCSC) transformation is accompanied by the breaking of old chemical bonds and the formation of new bonds in the crystalline state, which can affect the dimensionality of complexes.<sup>1</sup> However, maintaining crystallinity during SCSC transformations is often challenging, which may contribute to their rarity and unpredictable nature.<sup>2,3</sup> Heating and light irradiation are two important external stimuli used to mediate SCSC transformations.<sup>4,5</sup> Whilst there are abundant examples of SCSC transformations reported for main group elements,<sup>6</sup> transition metals,<sup>7</sup> and rare-earth metals,<sup>8</sup> instances of such transformations for actinide elements are few.<sup>9</sup> Based on our ongoing interest in uranium chemistry supported by double layer N-P ligands,<sup>10</sup> we hereby report the isolation of monomeric and dimeric uranium azides **2** and **3** through salt metathesis between **1** and  $NaN_3$  at different temperatures. A temperature-induced irreversible SCSC transformation from **3** to **2** was observed. In addition, the reactivity of **2** and **3** with  $KC_8$  was investigated, leading to the isolation of uranium nitride **4**.

Treatment of complex **1** with two equivalents of  $NaN_3$  in THF at room temperature (RT) for 10 h afforded a mixture

containing both yellow crystalline complex **2** and green crystalline complex **3** after recrystallization at RT (Scheme 1). Interestingly, upon storing the filtrate of this reaction at  $60^\circ C$ , we observed that the color of the filtrate changed from green to yellow, and the crystals of **2** were isolated with a 78% yield. Conversely, when the filtrate of this reaction was stored at  $-20^\circ C$  overnight, the crystals of **3** were formed with a 76% isolated yield (Scheme 1).

The structures of complexes **2** and **3** have been confirmed by single-crystal X-ray diffraction. Complex **2** is a monomer containing a single U center, which is coordinated by two mono-anionic N-P ligands and two  $N_3^-$  units (Fig. 1). The bond lengths of U1–N1/N1' (2.6990(18) Å) are longer than those of U1–N2/N2' (2.2665(18) Å), reflecting the dative bond between U1 and N1/N1'. The bond lengths of U1–N3, N3–N4, and N4–N5 are 2.340(2) Å, 1.157(3) Å, and 1.152(3) Å, respectively. These bond lengths are consistent with the structure of  $U-N=N^+=N^-$  for the terminal azide unit. The bond angles of U1–N3–N4 and N3–N4–N5 are  $159.0(2)^\circ$  and  $179.7(3)^\circ$ , respectively.

Complex **3** is a dimer containing two U centers connected by two  $N_3^-$  units (Fig. 2). The bond angles of U1–N8–N9 ( $140.0(10)^\circ$ ) and U1–N10–N11 ( $142.0(9)^\circ$ ) are smaller than the U1–N3–N4 angle ( $159.0(2)^\circ$ ) in complex **2**. The bond lengths of U1–N8 (2.546(11) Å) and U1–N10 (2.484(10) Å) are longer than the bond



Scheme 1 Synthesis of complexes **2**–**4**.

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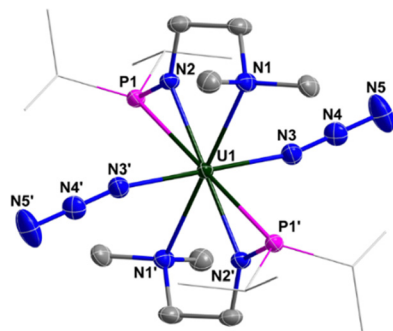


Fig. 1 Molecular structure of **2** with thermal ellipsoids at the 50% probability level. Hydrogen atoms are omitted for clarity. The <sup>i</sup>Pr groups on P atoms were simplified into lines.

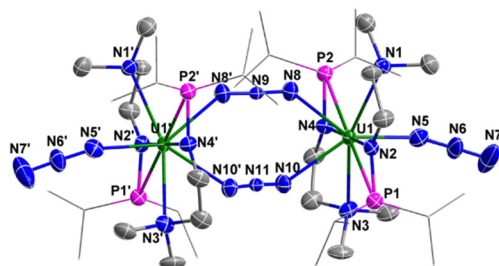


Fig. 2 Molecular structure of **3** with thermal ellipsoids at the 50% probability level. Hydrogen atoms are omitted for clarity. The <sup>i</sup>Pr groups on P atoms were simplified into lines.

length of U1–N5 (2.331(8) Å) due to the bridging nature of N8 and N10. Correspondingly, the bond lengths of N8–N9 (1.164(11) Å) and N10–N11 (1.180(10) Å) in the bridged N<sub>3</sub><sup>−</sup> units are longer than those in the terminal N<sub>3</sub><sup>−</sup> units, such as N6–N7 (1.151(14) Å).

In solution, complexes **2** and **3** can be interconverted at −20 °C and 60 °C. Consistently, the <sup>1</sup>H NMR spectra of complexes **2** or **3** recorded at RT demonstrated a 1:1 ratio between the monomer and dimer forms (Fig. S5 and S6, ESI†). Therefore, variable-temperature <sup>1</sup>H NMR spectra of complex **3** were measured in d<sub>8</sub>-THF from 213 K to 333 K (Fig. 3). At 213 K, a range of signals between +82.1 and −76.3 ppm was attributed to resonances for complex **3**. As the temperature increased, the signals for complex **3** gradually disappeared, and a set of broad signals for complex **2** emerged, indicating the conversion from **3** to **2**. From these variable-temperature <sup>1</sup>H NMR spectral data, we calculated Δ*H* to be 3.2 ± 0.4 kcal mol<sup>−1</sup> (Fig. S14, ESI†), which is in close proximity to the value obtained from DFT calculations (5.0 kcal mol<sup>−1</sup>, Fig. S18, ESI†). Upon lowering the temperature from 333 K to 213 K, the signals for **2** disappeared and those for complex **3** reappeared (Fig. S1, ESI†). These results suggest that temperature-induced invertible transformation between complexes **2** and **3** is feasible in solution. The Fourier-transform infrared (FT-IR) spectra of complexes **2** and **3** were measured in the solid state at room temperature (Fig. S10 and S11, ESI†). Complex **3** exhibits two strong azide stretching vibrations at 2079 and 2134 cm<sup>−1</sup>, characteristic of U-bound terminal and bridged azide ligands, whereas complex **2** shows a

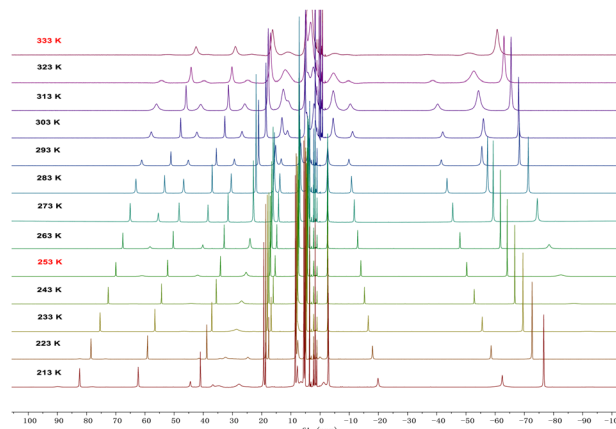


Fig. 3 Variable-temperature <sup>1</sup>H NMR spectra (from 213 K to 333 K) of complex **3** in d<sub>8</sub>-THF.

single strong vibration at 2075 cm<sup>−1</sup>, indicative of U-bound terminal azide ligands.<sup>10a,11</sup>

The UV-Vis-NIR electronic absorption spectra of complexes **2** and **3** were measured in THF at RT (Fig. S7, ESI†). Complexes **2** and **3** display an intense absorption peak at 227 nm and 238 nm, respectively. In addition, a weak absorption peak at 452 nm for complex **2** and at 456 nm for complex **3** is also observed in the visible region. Furthermore, a series of low-intensity absorptions are observed in the NIR region. For example, peaks at 821, 893, 1129, 1339, 1409, and 1510 nm were observed for complex **2**, while peaks at 819, 1129, 1335, 1427, and 1525 nm were observed for complex **3**. These low-intensity absorptions ( $\epsilon < 200 \text{ M}^{-1} \text{ cm}^{-1}$ ) in the NIR region for complexes **2** and **3** are attributed to the 5*f*–5*f* transitions, which are typical characteristics of U(IV) species.

Variable-temperature magnetic data for complexes **2** and **3** in the solid state were collected using a superconducting quantum interference device (SQUID) magnetometer (Fig. 4). The magnetic moment of complex **2** at 370 K is 3.10 μ<sub>B</sub>, which declines slowly to 2.49 μ<sub>B</sub> at 50 K, and then drops sharply to 0.55 μ<sub>B</sub> at 1.8 K with a trend toward zero. The effective moment for complex **3** at 1.8 K is 0.87 μ<sub>B</sub>, which increases sharply to 3.02 μ<sub>B</sub> at 50 K. With an increase in temperature from 50 K to 340 K, the magnetic moment of complex **3** reaches 3.31 μ<sub>B</sub>.

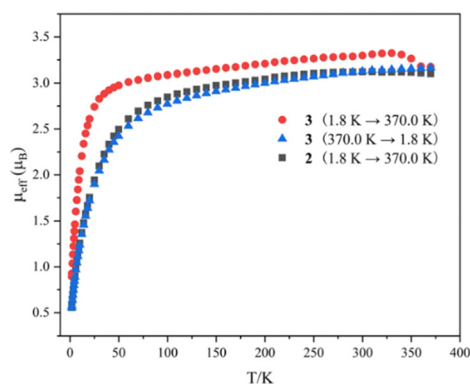


Fig. 4 Variable-temperature effective magnetic moment data. Magnetic moment per uranium ion for **2** and **3**.



When the temperature rises from 340 K to 370 K, the magnetic moments of complex **3** drop noticeably from 3.31 to 3.16  $\mu_B$ , which is comparable to the moment of complex **2** at 370 K (3.10  $\mu_B$ ). With a reduction in temperature from 370 K to 1.8 K, the magnetic moment of complex **3** exhibits a similar trend and values to those of complex **2**. The magnitudes of  $\mu_{\text{eff}}$  and the temperature dependence for complexes **2** and **3** are comparable to those observed for previously reported U(IV) complexes.<sup>12</sup> These results demonstrate that the dimeric complex **3** converts to the monomeric complex **2** at around 350 K in the solid state, suggesting a probable SCSC process.

To corroborate the process of SCSC transformation, two approaches were employed. Initially, a crystal of complex **3** was placed on a diffractometer and X-ray diffraction data were collected at different temperatures (313 K, 323 K, 333 K, 343 K, and 353 K). Analysis of these data revealed that complex **3** underwent a transformation to complex **2** at 353 K (Table S2, ESI<sup>†</sup>), which is further corroborated by a color change of the crystal from green to yellow at this temperature (Fig. S15, ESI<sup>†</sup>). The second method involved heating a crystal of complex **3** at incremental temperatures (40 °C, 50 °C, 60 °C, 70 °C, and 80 °C) within a glovebox for 20 minutes, followed by X-ray diffraction data collection at 193 K. The analysis of these data also confirmed that complex **3** converted to complex **2** after heating at 80 °C (Table S3, ESI<sup>†</sup>). Photographic evidence of the crystal additionally supports this transformation (Fig. S16, ESI<sup>†</sup>). The persistence of crystallinity during heating suggests an authentic SCSC transformation process.

Calculations were performed on **2** and **3** at the DFT level using the B3PW91 functional and including solvent effects (see ESI<sup>†</sup> for details). The formation of complex **2** from precursor **1** is exothermic by 16.5 kcal mol<sup>−1</sup>. The dimerization of complex **2** to yield complex **3** is found to be exothermic by 5.0 kcal mol<sup>−1</sup>, in line with the thermal equilibrium between complexes **2** and **3** reported experimentally. The optimized geometry is agreement with the experimental one, particularly the unsymmetrical coordination of the monoanionic N–P ligand (with U–N1 and U–N2 distances of 2.27 Å and 2.71 Å, respectively). Similarly, the azide coordination to the uranium is well reproduced, with the U–N3 distance being 2.35 Å. Natural Bonding Orbital (NBO) analysis indicates the presence of a polarized U–N3 bond (90% character on N) with a Wiberg Bond Index (WBI) of 0.78. The N3–N4 and N4–N5 WBIs are 1.71 and 2.14, respectively, consistent with the presence of two relatively covalent N=N double bonds interacting with a U(IV) center.

Geometry optimization was also performed on complex **3**, and different spin states were considered, namely the quintet, triplet, and open-shell singlet. The quintet and open-shell singlet states were found to be degenerate, while the triplet state is 10.5 kcal mol<sup>−1</sup> higher in energy. The unpaired spin density plots (Fig. S20, ESI<sup>†</sup>) show that both uranium centers are in a +IV oxidation state. The distinction between the quintet and the open-shell singlet states lies in the spin coupling between the two U(IV) centers—ferromagnetic for the quintet and anti-ferromagnetic for the open-shell singlet—with the small energy difference indicating that there is no strong coupling between

them. The associated UV-visible spectrum (Fig. S22, ESI<sup>†</sup>), computed in THF solution, displays intense absorption in the UV region with an absorption feature in the visible region (around 400 nm), corroborating the experimental observations.

Uranium azides are well-known precursors for the synthesis of uranium nitride *via* photolysis or reduction. However, the photolysis of complex **2** under a 40 W UV lamp for 10 hours only generates intractable results (Fig. S9, ESI<sup>†</sup>). Fortunately, treating complex **2** with two equivalents of KC<sub>8</sub> in THF affords a brownish solid, complex **4**, in a 67% yield after recrystallization from Et<sub>2</sub>O at RT (Scheme 1). Complex **4** can also be synthesized by reducing complex **3** with four equivalents of KC<sub>8</sub> under the same conditions. The <sup>1</sup>H NMR spectrum of **4** exhibits five signals ranging from +7.6 to −30.1 ppm, with the signal of the N–CH<sub>3</sub> group located at −30.1 ppm.

The solid-state structure of complex **4** was determined by single-crystal X-ray diffraction (Fig. 5). The diamond-shaped U<sub>2</sub>N<sub>2</sub> core is supported by four monoanionic N–P ligands. The U1–N9 (2.090(9) Å), U1–N10 (2.113(7) Å), U2–N9 (2.087(7) Å) and U2–N10 (2.068(8) Å) bond lengths are comparable to the U–N bond lengths in [K(DME)(calix[4]tetrapyrrole)U<sup>V</sup>]<sub>2</sub>(μ-NK)<sub>2</sub>[K(DME)<sub>4</sub>] (2.076(6) and 2.099(5) Å),<sup>13</sup> [KU<sup>V</sup>(μ-N)(OSi(O<sup>t</sup>Bu)<sub>3</sub>)<sub>3</sub>]<sub>2</sub> (2.022(5) and 2.101(6) Å),<sup>14</sup> [Cs<sub>3</sub>{U<sup>V</sup>(OSi(O<sup>t</sup>Bu)<sub>3</sub>)<sub>3</sub>}(μ-N)<sub>2</sub>U<sup>V</sup>(OSi(O<sup>t</sup>Bu)<sub>3</sub>)<sub>2</sub>(κ-O)] [CsOSi(O<sup>t</sup>Bu)<sub>3</sub>] (1.85(1), 2.09(1), 2.13(1) and 2.34(1) Å),<sup>15</sup> [(μ-N)(U<sup>V</sup>(N<sup>t</sup>Bu)Ar)<sub>3</sub>]<sub>2</sub>[B(Ar<sup>F</sup>)<sub>4</sub>] (Ar = 3,5-Me<sub>2</sub>C<sub>6</sub>H<sub>3</sub>, Ar<sup>F</sup> = 3,5-(CF<sub>3</sub>)<sub>2</sub>C<sub>6</sub>H<sub>3</sub>) (2.0470(3) and 2.0511(3) Å),<sup>16</sup> and in U(IV) nitride containing U<sub>2</sub>N<sub>2</sub> core (2.153(2) Å).<sup>17</sup> However, these U–N bond lengths are slightly longer than terminal uranium(v) nitrides supported by Tren<sup>TIPS</sup> ligand (1.80–1.89 Å).<sup>18</sup>

The UV-visible-NIR electronic absorption spectrum of complex **4** was measured in THF at room temperature (Fig. S12, ESI<sup>†</sup>), displaying an intense absorption peak at 225 nm and a series of low-intensity absorptions in the NIR region. These broad features in the NIR region are comparable to those reported for U(v) nitride.<sup>11c,14,19</sup> The effective magnetic moment at 300 K is 2.94  $\mu_B$  (Fig. S13, ESI<sup>†</sup>), which is close to the theoretical value of 2.54  $\mu_B$  for a U(v) ion.<sup>10f,12</sup> With decreasing temperature, the moment decreases to 0.83  $\mu_B$  at 1.8 K per uranium. The variation in  $\mu_{\text{eff}}$  and the temperature dependence

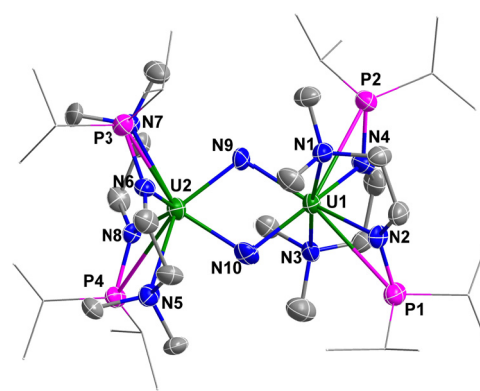


Fig. 5 Molecular structure of **4** with thermal ellipsoids at the 50% probability level. Hydrogen atoms are omitted for clarity. The 'Pr groups on P atoms were simplified into lines.



of complex **4** are comparable to those of reported U(v) complexes.<sup>20</sup>

Finally, calculations on complex **4** considered three different spin states. The triplet and open-shell singlet were found to be degenerate, whereas the closed-shell singlet is higher in energy by 30.7 kcal mol<sup>-1</sup>. The weak coupling between the two U centers is indicated by the small energy difference between the ferro- and antiferromagnetic states. The U=N Wiberg Bond Indices (WBIs) are 1.43 and 1.37, consistent with the presence of delocalized U=N double bonds. The U-U WBI is 0.5, highlighting electron delocalization within the U<sub>2</sub>N<sub>2</sub> moiety.

In summary, monomeric and dimeric uranium azides, complexes **2** and **3**, were isolated by reacting complex **1** with NaN<sub>3</sub> at different temperatures. A temperature-induced irreversible SCSC transformation was observed from complex **3** to complex **2**. The uranium nitride complex **4**, featuring a U<sub>2</sub>N<sub>2</sub> core, was isolated by reducing complexes **2** or **3** with KC<sub>8</sub>.

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## Conflicts of interest

The authors declare no competing financial interests.

## Notes and references

- (a) G. Chakraborty, I.-H. Park, R. Medishetty and J. J. Vittal, *Chem. Rev.*, 2021, **121**, 3751; (b) H. Wang, W. Meng, J. Wu, J. Ding, H. Hou and Y. Fan, *Coord. Chem. Rev.*, 2016, **307**, 130; (c) S. Kitagawa and K. Uemura, *Chem. Soc. Rev.*, 2005, **34**, 109; (d) Y. Chen, Q. Chen and Z. Zhang, *Chin. J. Org. Chem.*, 2021, **41**, 3826.
- (a) J. J. Vittal, *Coord. Chem. Rev.*, 2007, **251**, 1781; (b) O. V. Zenkina, E. C. Keske, R. Wang and C. M. Crudden, *Angew. Chem., Int. Ed.*, 2011, **50**, 8100; (c) S. Rodríguez-Jiménez, H. L. C. Feltham and S. Brooker, *Angew. Chem., Int. Ed.*, 2016, **55**, 15067.
- (a) C. G. Bezzu, M. Helliwell, J. E. Warren, D. R. Allan and N. B. Mckeown, *Science*, 2010, **327**, 1627; (b) M. Kawano and M. Fujita, *Coord. Chem. Rev.*, 2007, **251**, 2592; (c) Q. Liu, M. Lu, L. Yu, Y. Song and J. Lang, *Chin. J. Chem.*, 2021, **39**, 647; (d) Y. Cui and D. Xia, *Chin. J. Org. Chem.*, 2021, **41**, 907.
- (a) G. K. Kole and J. J. Vittal, *Chem. Soc. Rev.*, 2013, **42**, 1755; (b) E. Fernandez-Bartolome, A. Martinez-Martinez, E. Resines-Urien, L. Pineiro-Lopez and J. S. Costa, *Coord. Chem. Rev.*, 2022, **452**, 214281; (c) W. He, S. Li and Y. Lan, *Inorg. Chem. Front.*, 2018, **5**, 279.
- (a) S. Huang, T. Andy Hor and G. Jin, *Coord. Chem. Rev.*, 2017, **346**, 112; (b) B. B. Rath and J. J. Vittal, *Acc. Chem. Res.*, 2022, **55**, 1445; (c) Akhtaruzzaman, S. Khan, B. Dutta, T. S. Kannan, G. K. Kole and M. H. Mir, *Coord. Chem. Rev.*, 2023, **483**, 215095; (d) M. Zhang, X. Yang, Z. Xu, J. Yin and S. Liu, *Chin. J. Chem.*, 2022, **40**, 2853.
- (a) A. J. Thompson, A. I. C. Orúe, A. J. Nair, J. R. Piec, J. McMurtrie and J. K. Clegg, *Chem. Soc. Rev.*, 2021, **50**, 11725; (b) A. K. Saini, K. Natarajan and S. M. Mobin, *Chem. Commun.*, 2017, **53**, 9870.
- (a) H. Li, S. Zhao, S. Zang and J. Li, *Chem. Soc. Rev.*, 2020, **49**, 6364; (b) E. A. Patrick, M. E. Bowden, J. D. Erickson, R. M. Bullock and B. L. Tran, *Angew. Chem., Int. Ed.*, 2023, e202304648.
- (a) Y. So and W. Leung, *Coord. Chem. Rev.*, 2017, **340**, 172; (b) W. Chen, Y. Chen, G. Huang, J. Liu, J. Jia and M. Tong, *Chem. Commun.*, 2018, **54**, 10886; (c) Y. Xin, J. Wang, M. Zychowicz, J. J. Zakrzewski, K. Nakabayashi, B. Sieklucka, S. Chorazy and S.-I. Ohkoshi, *J. Am. Chem. Soc.*, 2019, **141**, 18211.
- (a) K. Lv, S. Fichter, M. Gu, J. Marz and M. Schmidt, *Coord. Chem. Rev.*, 2021, **446**, 214011; (b) T. Zheng, Y. Gao, D. Gui, L. Chen, D. Sheng, J. Diwu, Z. Chai, T. Albrecht-Schmitt and S. Wang, *Dalton Trans.*, 2016, **45**, 9031; (c) Z. Xie, L. Mei, Q. Wu, K. Hu, L. Xia, Z. Chai and W. Shi, *Dalton Trans.*, 2017, **46**, 7392; (d) L. Chen, Y. Zhang, Z. Weng, Z. Liu, J. Zhang, Y. Wang and S. Wang, *Chin. J. Chem.*, 2021, **39**, 597; (e) L. Mei, L. Wang, L. Yuan, S. An, Y. Zhao, Z. Chai, P. C. Burns and W. Shi, *Chem. Commun.*, 2015, **51**, 11990.
- (a) X. Xin, I. Douair, Y. Zhao, S. Wang, L. Maron and C. Zhu, *Natl. Sci. Rev.*, 2023, **10**, nwac144; (b) X. Xin, I. Douair, T. Rajeshkumar, Y. Zhao, S. Wang, L. Maron and C. Zhu, *Nat. Commun.*, 2022, **13**, 3809; (c) X. Xin, I. Douair, Y. Zhao, S. Wang, L. Maron and C. Zhu, *J. Am. Chem. Soc.*, 2020, **142**, 15004; (d) P. Wang, Y. Zhao and C. Zhu, *Organometallics*, 2022, **41**, 2448; (e) P. Wang, I. Douair, Y. Zhao, S. Wang, J. Zhu, L. Maron and C. Zhu, *Angew. Chem., Int. Ed.*, 2021, **60**, 473; (f) J. Shen, T. Rajeshkumar, G. Feng, Y. Zhao, S. Wang, L. Maron and C. Zhu, *Angew. Chem., Int. Ed.*, 2023, **62**, e202303379; (g) G. Feng, M. Zhang, D. Shao, X. Wang, S. Wang, L. Maron and C. Zhu, *Nat. Chem.*, 2019, **11**, 248; (h) G. Feng, M. Zhang, P. Wang, S. Wang, L. Maron and C. Zhu, *Proc. Natl. Acad. Sci. U. S. A.*, 2019, **116**, 17654; (i) P. Wang, I. Douair, Y. Zhao, R. Ge, J. Wang, S. Wang, L. Maron and C. Zhu, *Chemistry*, 2022, **8**, 1361; (j) G. Feng, K. N. McCabe, S. Wang, L. Maron and C. Zhu, *Chem. Sci.*, 2020, **11**, 7585; (k) W. Fang, S. Pan, W. Su, S. Wang, L. Zhao, G. Frenking and C. Zhu, *CCS Chem.*, 2022, **4**, 1921; (l) Q. Zhu, W. Fang, L. Maron and C. Zhu, *Acc. Chem. Res.*, 2022, **55**, 1718; (m) W. Fang, Q. Zhu and C. Zhu, *Chem. Soc. Rev.*, 2022, **51**, 8434; (n) X. Sun, X. Gong, Z. Xie and C. Zhu, *Chin. J. Chem.*, 2022, **40**, 2047; (o) W. Fang, I. Douair, A. Hauser, K. Li, Y. Zhao, P. W. Roesky, S. Wang, L. Maron and C. Zhu, *CCS Chem.*, 2022, **4**, 2630; (p) K. Li, J. He, Y. Zhao and C. Zhu, *Inorg. Chem. Front.*, 2023, **10**, 5622.
- (a) R. K. Thomson, T. Cantat, B. L. Scott, D. E. Morris, E. R. Batista and J. L. Kiplinger, *Nat. Chem.*, 2010, **2**, 723; (b) R. Haiges, M. Vasiliu, D. A. Dixon and K. O. Christe, *Chem. – Eur. J.*, 2017, **23**, 652; (c) J. Du, D. M. King, L. Chatelain, E. Lu, F. Tuna, E. J. L. McInnes, A. J. Woolees, L. Maron and S. T. Liddle, *Chem. Sci.*, 2019, **10**, 3738.
- D. R. Kindra and W. J. Evans, *Chem. Rev.*, 2014, **114**, 8865.
- I. Korobkov, S. Gambarotta and G. Yap, *Angew. Chem., Int. Ed.*, 2002, **41**, 3433.
- (a) C. Camp, J. Pécaut and M. Mazzanti, *J. Am. Chem. Soc.*, 2013, **135**, 12101; (b) L. Barluzzi, L. Chatelain, F. Fadaei-Tirani, I. Zivkovic and M. Mazzanti, *Chem. Sci.*, 2019, **10**, 3543; (c) L. Barluzzi, F.-C. Hsueh, R. Scopelliti, B. E. Atkinson, N. Kaltsayannis and M. Mazzanti, *Chem. Sci.*, 2021, **12**, 8096.
- N. Jori, T. Rajeshkumar, R. Scopelliti, I. Zivkovic, A. Sienkiewicz, L. Maron and M. Mazzanti, *Chem. Sci.*, 2022, **13**, 9232.
- A. R. Fox, P. L. Arnold and C. C. Cummins, *J. Am. Chem. Soc.*, 2010, **132**, 3250.
- D. M. King, B. E. Atkinson, L. Chatelain, M. Gregson, J. A. Seed, A. J. Woolees, N. Kaltsayannis and S. T. Liddle, *Dalton Trans.*, 2022, **51**, 8855.
- (a) D. M. King, F. Tuna, E. J. L. McInnes, J. McMaster, W. Lewis, A. J. Blake and S. T. Liddle, *Nat. Chem.*, 2013, **5**, 482; (b) D. M. King, P. A. Cleaves, A. J. Woolees, B. M. Gardner, N. F. Chilton, F. Tuna, W. Lewis, E. J. L. McInnes and S. T. Liddle, *Nat. Commun.*, 2016, **7**, 13773.
- P. A. Cleaves, D. M. King, C. E. Kefalidis, L. Maron, F. Tuna, E. J. L. McInnes, J. McMaster, W. Lewis, A. J. Blake and S. T. Liddle, *Angew. Chem., Int. Ed.*, 2014, **53**, 10412.
- (a) P. L. Arnold, A.-F. Pecharman, E. Hollis, A. Yahia, L. Maron, S. Parsons and J. B. Love, *Nat. Chem.*, 2010, **2**, 1056; (b) O. P. Lam, F. W. Heinemann and K. Meyer, *C. R. Chim.*, 2010, **13**, 803; (c) D. Patel, F. Moro, J. McMaster, W. Lewis, A. J. Blake and S. T. Liddle, *Angew. Chem., Int. Ed.*, 2011, **50**, 10388; (d) O. P. Lam, F. W. Heinemann and K. Meyer, *Chem. Sci.*, 2011, **2**, 1538; (e) D. M. King, F. Tuna, E. J. L. McInnes, J. McMaster, W. Lewis, A. J. Blake and S. T. Liddle, *Science*, 2012, **337**, 717; (f) D. Patel, F. Tuna, E. J. L. McInnes, J. McMaster, W. Lewis, A. J. Blake and S. T. Liddle, *Dalton Trans.*, 2013, **42**, 5224; (g) C. T. Palumbo, L. Barluzzi, R. Scopelliti, I. Zivkovic, A. Fabrizio, C. Corminboeuf and M. Mazzanti, *Chem. Sci.*, 2019, **10**, 8840.

