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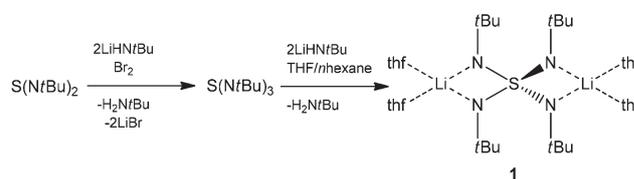
## Transition metal complexes containing the $S(NtBu)_4^{2-}$ tetraimidosulfate dianion†‡

Julia Matussek, Regine Herbst-Irmer, Ina Objartel and Dietmar Stalke\*

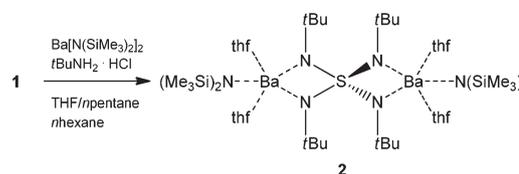
Three novel metal complexes [(*acac*)<sub>2</sub>Cu<sub>2</sub>(*NtBu*)<sub>4</sub>S] (**3**), [Li(*thf*)<sub>4</sub>]<sub>2</sub>[I<sub>4</sub>Cd<sub>2</sub>(*NtBu*)<sub>4</sub>S] (**4**) and [(*thf*)<sub>2</sub>Li(N(SiMe<sub>3</sub>)<sub>2</sub>N)-Zn(*NtBu*)<sub>4</sub>S] (**5**) are prepared from the intended transmetalation of the dilithium complex of the dilithium complex of *N,N,N',N''*-tetrakis(*tert*-butyl)tetraimidosulfate [(*thf*)<sub>4</sub>Li<sub>2</sub>(*NtBu*)<sub>4</sub>S] (**1**). The two lithium cations are replaced by either the cationic (*acac*)Cu(II) moiety, the neutral I<sub>2</sub>Cd(II) residue or only a single lithium cation is substituted by the cationic (Me<sub>3</sub>Si)<sub>2</sub>NZn(II) fragment. The complexes show two main results: first the S(*NtBu*)<sub>4</sub><sup>2-</sup> tetrahedron can serve as a ligand to transition metals from the soft Cu(II) to the harder Zn(II) at opposite sides and second the S–N bond distances vary only marginally in response to the various metals and the four distances constantly sum up to 6.38(2) Å. Hence the electropositive sulfur atom responds by internal shift to the metal-polarized negative charge at the outside of the S(NR)<sub>4</sub><sup>2-</sup> tetrahedron.

Isovalent electronic replacement of the oxygen atoms in the classic SO<sub>*n*</sub><sup>*m*-</sup> molecules and ions by NR imido groups yields the polyimido sulfur species S(NR)<sub>*n*</sub><sup>*m*-</sup> (*n* = 2, 3, 4 and *m* = 0, 2).<sup>1–8</sup> By introducing organic substituents to the chelating nitrogen atoms, the polyanion becomes more lipophilic, thus, the resulting complexes are frequently soluble in non-polar hydrocarbons and stay in the molecular regime rather than aggregate like their S–O counterparts.<sup>6</sup> Due to the large variety of coordination modes as found for sulfate anions, these polyimido compounds hold interesting electronic and stereochemical properties.<sup>9–14</sup> In 1997 we first synthesized the starting material to the current paper, dilithium-*N,N,N',N''*-tetrakis(*tert*-butyl)tetraimido sulfate (**1**) (Scheme 1) in a dual addition reaction of first lithium amide to S(*NtBu*)<sub>2</sub> to give the product S(*NtBu*)<sub>3</sub> upon oxidation with bromine.<sup>15,16</sup> Subsequently another equivalent of lithium *tert*-butylamide is added to the sulfur triimide to give **1** (Scheme 1). Once the S(NR)<sub>4</sub><sup>2-</sup> scaffold was synthesized in the following year the barium complex [(*thf*)<sub>4</sub>Ba<sub>2</sub>{N(SiMe<sub>3</sub>)<sub>2</sub>}{(*NtBu*)<sub>4</sub>S}] (**2**) could be obtained (Scheme 2) by first protonating **1** with *tert*-butylammonium chloride to give (*t*BuNH)<sub>2</sub>S(*NtBu*)<sub>2</sub> and subsequent metallation with [Ba{N(SiMe<sub>3</sub>)<sub>2</sub>}<sub>2</sub>].<sup>17</sup>

Thus, these previous results indicate that S(NR)<sub>4</sub><sup>2-</sup> can be coordinated by main group metals such as lithium and



**Scheme 1** Preparation of dilithium-*N,N,N',N''*-tetrakis(*tert*-butyl)tetraimidosulfate (**1**).<sup>15,16</sup>



**Scheme 2** Preparation of [(*thf*)<sub>4</sub>Ba<sub>2</sub>{N(SiMe<sub>3</sub>)<sub>2</sub>}{(*NtBu*)<sub>4</sub>S}] (**2**).<sup>17</sup>

barium, but the coordination to transition metals remained unknown. Hence we now embarked to synthesize d-block metal complexes, selecting late transition metals first as they resemble alkaline and alkaline earth metal properties best. By the preparation and isolation of [(*acac*)<sub>2</sub>Cu<sub>2</sub>(*NtBu*)<sub>4</sub>S] (**3**), [Li(*thf*)<sub>4</sub>]<sub>2</sub>[I<sub>4</sub>Cd<sub>2</sub>(*NtBu*)<sub>4</sub>S] (**4**) and [(*thf*)<sub>2</sub>Li(N(SiMe<sub>3</sub>)<sub>2</sub>)Zn(*NtBu*)<sub>4</sub>S] (**5**) presented herein we show for the first time that the lithium cations in **1** can be replaced by the transition metals copper(II), zinc(II) and cadmium(II). Apart from their similarity to s-block metals these metals were picked because of their divalent character, availability and potential application in catalysis. They will considerably widen the scope of the established sulfur imido chemistry.<sup>8,18</sup>

Institut für Anorganische Chemie der Georg-August-Universität Göttingen,  
Tammannstr. 4, 37077 Göttingen, Germany.

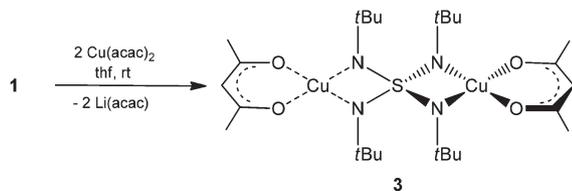
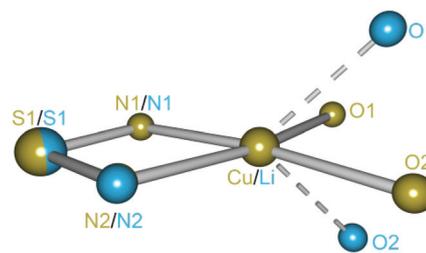
E-mail: dstalke@chemie.uni-goettingen.de; Fax: +49 551 39 33373;

Tel: +49 551 39 33000

† In memory of Professor Ken Wade.

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Scheme 3 Synthesis of  $[(\text{acac})_2\text{Cu}_2(\text{NtBu})_4\text{S}]$  (**3**).Fig. 2 Superposition plot of **1** (Li, light blue) and **3** (Cu, brass). The atoms S1, N1 and N2 are projected onto each other with a deviation of 0.0161 Å.

## Results and discussion

### Synthetic and structural studies

The copper complex  $[(\text{acac})_2\text{Cu}_2(\text{NtBu})_4\text{S}]$  (**3**) can be isolated from the reaction of copper acetylacetonate with  $[(\text{thf})_4\text{Li}_2(\text{NtBu})_4\text{S}]$  (**1**) according to Scheme 3. Two equivalents of lithium acetylacetonate precipitate and are removed by filtration. After one week in THF at  $-24\text{ }^\circ\text{C}$  colorless blocks, suitable for X-ray structure determination, were obtained at a yield of 59%. **3** crystallizes in the monoclinic space group  $P2_1/n$  with half of the molecule and one THF molecule in the asymmetric unit.

Each copper(II) atom is fourfold coordinated by the two oxygen atoms of the planar chelating acetylacetonate anion and by two nitrogen atoms of two opposite sides of the  $\text{S}(\text{NtBu})_4^{2-}$  tetrahedron. The fourfold coordination at the  $\text{Cu}(\text{II})$  atom can be described as a nearly square planar environment ( $\text{O1}-\text{Cu1}-\text{N1}$ :  $169.79^\circ$ ,  $\text{O2}-\text{Cu1}-\text{N2}$ :  $169.33^\circ$ ). This differs considerably from the nearly tetrahedral  $\text{N}_2\text{O}_2$ -coordination of the lithiated starting material **1** (Fig. 1 and 2).

The two crystallographically independent S–N bond lengths in **3** ( $1.59\text{ \AA}$ ) do not differ significantly from the S–N bond lengths in **1** ( $1.60\text{ \AA}$ ) and are half-way between the values normally quoted for a typical S–N single bond ( $1.69\text{ \AA}$ )<sup>19</sup> and a S–N double bond ( $1.52\text{ \AA}$ ).<sup>19</sup> However, for none of the investigated S–N bonds in methyl(diimido)sulfinic acid  $\text{H}(\text{NtBu})_2\text{SME}$

( $1.68$  and  $1.58\text{ \AA}$ ), methylene-bis(triimido)sulfonic acid  $\text{H}_2\text{C}\{\text{S}(\text{NtBu})_2(\text{NHtBu})\}_2$  ( $1.52$  to  $1.65\text{ \AA}$ ), sulfurdiimide  $\text{S}(\text{NtBu})_2$  ( $1.54$  and  $1.53\text{ \AA}$ ), and sulfurtriimide  $\text{S}(\text{NtBu})_3$  ( $1.51\text{ \AA}$ ), a classical double bond formulation could be supported from charge density investigations.<sup>20</sup> This was further substantiated by the NBO/NRT approach. Valence expansion to more than eight electrons at the sulfur atom can definitely be excluded to explain the bonding.<sup>17,21</sup> The same was shown recently for the sulfate anion,  $\text{SO}_4^{2-}$ , as well by charge density based both on experimental and theoretical methods.<sup>22</sup>

Due to the similar bond lengths, equal distribution of the two negative charges over the four nitrogen atoms of the  $\text{S}(\text{NR})_4^{2-}$ -ligand is assumed. Furthermore, the  $\text{N}\cdots\text{M}$  coordination (**3/1**:  $1.96\text{ \AA}$ ) and the angles  $\text{N}\cdots\text{M}\cdots\text{N}$  (**3**:  $72.29^\circ$ , **1**:  $73.95^\circ$ ) and  $\text{N}-\text{S}-\text{N}$  (**3**:  $93.47^\circ$ , **1**:  $94.60^\circ$ ) are comparable. This can be explained by the similar cationic radius of  $\text{Cu}(\text{II})$  and  $\text{Li}(\text{I})$  ( $\text{Cu}^{2+}$ :  $0.71\text{ \AA}$ ,  $\text{Li}^+$ :  $0.73\text{ \AA}$ ).<sup>23</sup> In published complexes with coordinated metal acetylacetonate at the nitrogen atom the  $\text{N}\cdots\text{M}$  distances are  $2.17\text{ \AA}$  on average but the  $\text{Cu}(\text{acac})^+$  cation in a fourfold coordination sphere attains  $\text{N}\cdots\text{M}$  distances of  $1.96\text{ \AA}$ <sup>24</sup> which is in excellent agreement with this result.

In the reaction of  $[(\text{thf})_4\text{Li}_2(\text{NtBu})_4\text{S}]$  (**1**) with cadmium iodide surprisingly  $[\text{Li}(\text{thf})_4]_2[\text{I}_4\text{Cd}_2(\text{NtBu})_4\text{S}]$  (**4**) is obtained according to Scheme 4. The anticipated transmetalation and salt elimination of  $\text{LiI}$ , which should be the driving force for the reaction, did not occur. Instead, the solvent separated ion pair  $[\text{Li}(\text{thf})_4]_2[\text{I}_4\text{Cd}_2(\text{NtBu})_4\text{S}]$  (**4**) is found, where two equivalents of cadmium(II) iodide are coordinated by two opposite sides of the  $\text{S}(\text{NtBu})_4^{2-}$  tetrahedron, resulting in the  $[\text{I}_4\text{Cd}_2(\text{NtBu})_4\text{S}]^{2-}$  dianion. Two tetrahedrally solvent coordinated  $[\text{Li}(\text{thf})_4]_2$  cations provide electro neutrality. This

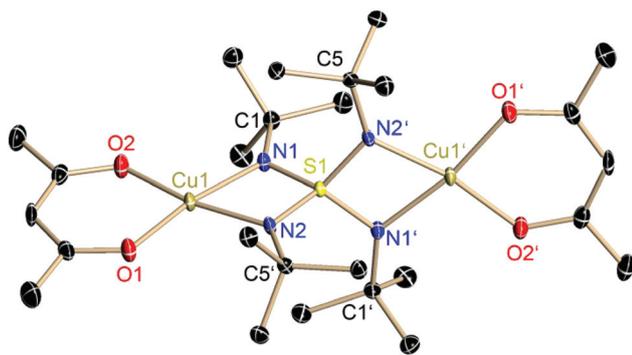
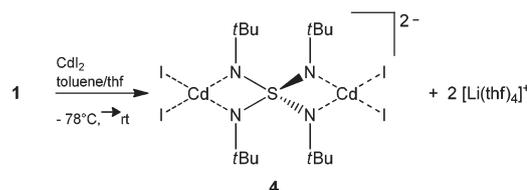


Fig. 1 Crystal structure of  $[(\text{acac})_2\text{Cu}_2(\text{NtBu})_4\text{S}]$  (**3**). Hydrogen atoms are omitted for clarity. Anisotropic displacement parameters are depicted at the 50% probability level. Selected bond lengths [ $\text{\AA}$ ] and angles [ $^\circ$ ]: S1–N1  $1.5867(14)$ , S1–N2  $1.547(14)$ , N1–Cu1  $1.9598(15)$ , N2–Cu1  $1.9555(15)$ , N1–S1–N2  $93.47(7)$ , N1–Cu1–N2  $72.29(6)$ .

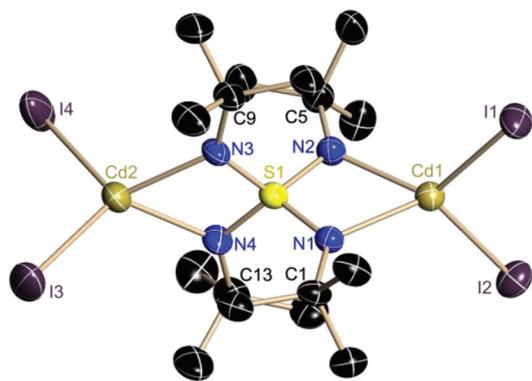
Scheme 4 Synthesis of  $[\text{Li}(\text{thf})_4]_2[\text{I}_4\text{Cd}_2(\text{NtBu})_4\text{S}]$  (**4**).

phenomenon might be explained by considering the various lattice energies of the involved metal halide salts. The lattice energy of CdI<sub>2</sub> with 2455 kJ mol<sup>-1</sup> (ref. 25) is overwhelmingly larger than the lattice energy of lithium iodide (2 × 746 kJ mol<sup>-1</sup> (ref. 25)). Thus, considering the energetic balance, elimination of LiI is unfavorable and a higher solvation enthalpy for the lithium ions with THF is accepted.

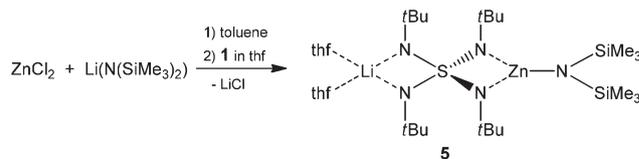
**4** crystallizes from THF–toluene within 4 weeks at –24 °C as colorless blocks in the monoclinic space group *P*<sub>2</sub><sub>1</sub>/*c* in a yield of 22%. The asymmetric unit contains one dianion, two cations and one THF molecule. The two cadmium atoms are tetrahedrally coordinated by two iodine atoms and two nitrogen atoms of the ligand. The central sulfur atom of the ligand possesses a distorted tetrahedral environment (95.3°–116.9°).

It is interesting to note that all three S–N bonds in all known metal complexes of the *S*-alkyltriimidatosulfonates [RS–(NR)<sub>3</sub>]<sup>–</sup> (M = Li, Ba, Al, Zn) and in the triimidatosulfonic acid MeS(NtBu)<sub>2</sub>NHtBu constantly sum up to 4.70(2) Å. The SN<sub>3</sub> unit responds flexibly to different electronic requirements induced by either different metal cations or conjugated *S*-substituents in terms of the sulfur atom being shifted relative to an otherwise fixed N<sub>3</sub> environment. This seems to be valid for the S(*ν*)–N bonds as well and experimentally emphasizes the predominantly ionic S–N bonding rather than valence expansion and *d*-orbital participation in bonding.<sup>26</sup> The four crystallographically independent S–N bond lengths average to 1.59 Å in **4** (Fig. 3).

While [(thf)<sub>4</sub>Ba<sub>2</sub>{N(SiMe<sub>3</sub>)<sub>2</sub>}{(NtBu)<sub>4</sub>S}] (**2**) is the product of a transmetalation using barium-hexamethylsilylamide from [(thf)<sub>4</sub>Li<sub>2</sub>(NtBu)<sub>4</sub>S] (**1**) via a hydrogenation first (Scheme 2)<sup>17</sup> we now report a transmetalation without the previous generation of a protonated species. Firstly, lithium hexamethylsilylamide and zinc chloride were reacted under elimination of lithium chloride to give the assumed intermediate Zn(Cl)N–



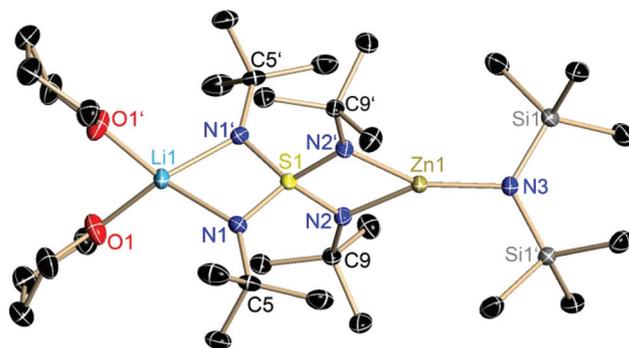
**Fig. 3** Crystal structure of the dianion in the solvent separated ion pair [Li(thf)<sub>4</sub>]<sub>2</sub>[Li<sub>4</sub>Cd<sub>2</sub>(NtBu)<sub>4</sub>S][thf] (**4**). Hydrogen atoms are omitted for clarity. Anisotropic displacement parameters are depicted at the 50% probability level. Selected bond lengths [Å] and angles [°]: S1–N1 1.585(5), S1–N2 1.595(5), S1–N3 1.592(5), S1–N4 1.589(5), N1–Cd1 2.227(5), N2–Cd1 2.215(5), N3–Cd2 2.227(5), N4–Cd2 2.217(5), Cd1–I1 2.7416(7), Cd1–I2 2.7489(7), Cd2–I3 2.7531(9), Cd2–I4 2.7372(8), N1–S1–N2 95.8(3), N3–S1–N4 95.2(3), N1–Cd1–N2 64.16(18), N3–Cd2–N4 63.81(18), I1–Cd1–I2 108.40(2), I3–Cd2–I4 109.34(3).



**Scheme 5** Synthesis of [(thf)<sub>2</sub>Li{(SiMe<sub>3</sub>)<sub>2</sub>N}Zn(NtBu)<sub>4</sub>S] (**5**).

(SiMe<sub>3</sub>). Subsequently, this intermediate gives the first isolated heterobimetallic compound [(thf)<sub>2</sub>Li{(SiMe<sub>3</sub>)<sub>2</sub>N}Zn(NtBu)<sub>4</sub>S] (**5**) after addition of **1** (Scheme 5).

**5** crystallizes from THF–toluene within four days at –24 °C, to give colorless blocks in a 34% yield, which were suitable for X-ray structure analysis. The compound crystallizes in the monoclinic space group *C*2/*c* with half a molecule per asymmetric unit. At one site of the S(NtBu)<sub>4</sub><sup>2–</sup> dianion a lithium ion remains coordinated like in the starting material and at the other site the zinc atom is complexed in a trigonal planar fashion by the two chelating nitrogen atoms of the ligand and on additional N(SiMe<sub>3</sub>)<sub>2</sub> amide group. In **5** the S1–N1 bond (1.5661(14) Å) is shorter than the S–N bond (1.59 Å) of the starting material because of the electron withdrawing effect of ZnN(SiMe<sub>3</sub>)<sub>2</sub><sup>+</sup>. While the harder Zn<sup>2+</sup> cation claims more negative charge from the two imide groups than the softer lithium cation the zinc-coordinated imide groups remain less attractive to the positively polarized sulfur atom (S1–N2 1.6312(14) Å), which compensates for its part at the lithium coordinated imide groups. The N1'–S1–N1 angle (96.80(11)) is wider than the N2'–S1–N2 angle (91.38(10)) presumably due to the higher steric demand of the (thf)<sub>2</sub>Li moiety compared to the N(SiMe<sub>3</sub>)<sub>2</sub> anion. The Li–N distance of 1.988(3) Å is typical for Li–N bonds.<sup>27</sup> Published distances between a lithium ion which is coordinated by two THF molecules and two nitrogen atoms, are on average 2.066 Å. The Zn–N(amide) distance is 1.880(2) Å which is only marginally shorter than the mean average of Zn–amide bonds in the CCDC (Fig. 4).<sup>28</sup>



**Fig. 4** Crystal structure of [(thf)<sub>2</sub>Li{(SiMe<sub>3</sub>)<sub>2</sub>N}Zn(NtBu)<sub>4</sub>S] (**5**). Hydrogen atoms are omitted for clarity. Anisotropic displacement parameters are depicted at the 50% probability level. Selected bond lengths [Å] and angles [°]: Li1–N1 1.988(3), N1–S1 1.5661(14), N2–S1 1.6312(14), N2–Zn1 1.9580(14), N3–Zn1 1.880(2), N2–S1–N2' 91.38(10), N1–S1–N1' 96.80(11), N1–Li1–N1' 72.18(15), S1–N1–Li1 95.51(10), S1–N2–Zn1 97.71(7), N2–Zn1–N3 143.41(4), N2–Zn1–N2' 73.19(8).



## Structural comparison

From the comparison of the three compounds it is interesting to note that in **2** the Cd–N bonds are longer than the M–N bonds in the other complexes, even longer than anticipated by the higher radius. They are further widened due to the lower electrostatic interactions to the neutral CdI<sub>2</sub> moieties compared to the otherwise cationic parts. Nevertheless, the complex to be formed indicates a certain binding affinity of the S(NR)<sub>4</sub><sup>2-</sup> ligand even to neutral residues.

As in the metal *S*-alkyltriamidosulfonates RS(NR)<sub>3</sub><sup>-</sup> also in the metal tetraimidosulfates S(NR)<sub>4</sub><sup>2-</sup> the sum of the S–N bond distances seems to be almost invariant to the metal coordination (4.70(2) in the first and 6.38(2) Å in the latter). In the rigid framework of the four electron-rich imido nitrogen atoms the electropositive sulfur atom is pulled towards the nitrogen atoms coordinated by the Li<sup>+</sup> because they remain more attractive as the Li<sup>+</sup> loses the competition for imide nitrogen density against the Zn<sup>2+</sup> and there remains more density at the LiN<sub>2</sub> site of the SN<sub>4</sub> tetrahedron. Hence the sulfur atom inside the N<sub>4</sub>-cage responds to the metal-polarized negative charge at the outside of the S(NR)<sub>4</sub><sup>2-</sup> dianion. This again emphasises predominantly ionic S–N bonding, reminiscent to S–O bonding in sulfate.<sup>22,29</sup>

## Experimental section

### General procedure

All experiments were performed either in an inert gas atmosphere of purified dry argon with standard Schlenk techniques<sup>30,31</sup> or in an argon glove box. The glassware was dried at 130 °C, assembled hot and cooled under reduced pressure. All solvents were dried over appropriate alkali metals, distilled and degassed prior to use. All NMR spectra were either recorded on a Bruker Avance DPX 300 MHz or Bruker Avance DRX 500 MHz spectrometer using TMS (<sup>1</sup>H, <sup>13</sup>C and <sup>29</sup>Si) and LiCl (<sup>7</sup>Li) as external reference and the protons of the deuterated solvents as internal standard. The spectra were measured at room temperature if not stated otherwise. Elemental analyses (C, H, N and S) were carried out at the Mikroanalytisches Labor, Institut für Anorganische Chemie, Universität Göttingen.

**[(acac)<sub>2</sub>Cu<sub>2</sub>(NtBu)<sub>4</sub>S] (3).** Cu(acac)<sub>2</sub> (77 mg, 0.294 mmol, 2.0 eq.) and [(thf)<sub>4</sub>Li<sub>2</sub>(NtBu)<sub>4</sub>S] (100 mg, 0.147 mmol, 1.0 eq.) were dissolved in THF (10 mL) and stirred overnight at room temperature. After removing lithium acetylacetonate by filtration and storing the green solution for 3 month at –24 °C, colorless crystals were obtained. Yield: 56 mg, 0.087 mmol, 59%; Elemental analysis (found (calc.) [%]): C 44.23 (48.65), H 7.47 (7.85), N 6.76 (8.73), S 4.94 (5.00). This poor elemental analysis is due to the contamination of the sample with approximately 25% silicon join grease (at 0.09 ppm in the <sup>1</sup>H-NMR and at 1.35 ppm in the <sup>13</sup>C-NMR for (OSiMe<sub>2</sub>)<sub>n</sub>). Due to paramagnetism the NMR-signals are very broad. <sup>1</sup>H-NMR (400.130 MHz, THF-d<sub>8</sub>): δ = 5.72 (s, 36 H, CH<sub>3</sub>), 3.88 (s, 12 H, CH<sub>3</sub>), –16.74 (s, 2 H, CH) ppm. <sup>15</sup>N-NMR (30.432 MHz,

THF-d<sub>8</sub>): δ = –269.0 (N) ppm. *m/z* [%]: 640 [(acac)<sub>2</sub>Cu<sub>2</sub>(NtBu)<sub>4</sub>S], 50, 365 [Cu(NtBu)<sub>4</sub>S], 50, 336 [(acac)Cu(NtBu)<sub>2</sub>S], 54, 304 [(acac)Cu(NtBu)<sub>2</sub>], 14, 136 (Cu(acac)), 8, 57 (tBu), 54.

**[Li(thf)<sub>4</sub>]<sub>2</sub>[I<sub>4</sub>Cd<sub>2</sub>(NtBu)<sub>4</sub>S] (4).** To [(thf)<sub>4</sub>Li<sub>2</sub>(NtBu)<sub>4</sub>S] (400 mg, 0.589 mmol, 1.0 eq.) and cadmium iodide (282 mg, 0.770 mmol, 1.3 eq.) toluene (10 mL) was added at –78 °C and stirred at room temperature overnight. After a week at –24 °C, 3 mL of THF were added and the solution was stored again at –24 °C. Colorless crystals were obtained after 4 weeks. Yield: 226 mg, 132 mmol, 22%; Elemental analysis (found (calc.) [%]): C 33.35 (35.16), H 6.13 (6.15), N 3.65 (3.42), S 2.19 (1.96). <sup>1</sup>H-NMR (500 MHz, THF-d<sub>8</sub>): δ (ppm) = 3.64–3.58 (m, 36 H, THF), 1.80–1.75 (m, 36 H, THF), 1.30 (s, 36 H, tBu); <sup>7</sup>Li-NMR (500 MHz, THF-d<sub>8</sub>): δ (ppm) = –0.11 (s, 2 Li); <sup>13</sup>C-NMR (500 MHz, THF-d<sub>8</sub>): δ (ppm) = 67.2–65.8 (m, 18 C-THF), 53.5 (s, 4 CCH<sub>3</sub>), 32.0 (s, 12 CCH<sub>3</sub>), 24.8–23.77 (m, 18 C-THF); <sup>113</sup>Cd-NMR (500 MHz, THF-d<sub>8</sub>): δ = –387.1 (s, 2 Cd).

**[(thf)<sub>2</sub>Li(N(SiMe<sub>3</sub>)<sub>2</sub>)Zn(NtBu)<sub>4</sub>S] (5).** A slurry of ZnCl<sub>2</sub> (80 mg, 0.591 mmol, 2.0 eq.) and Li(N(SiMe<sub>3</sub>)<sub>2</sub>) (120 mg, 0.591 mmol, 2.0 eq.) in toluene (3 mL) was stirred 4 h at room temperature. To the white solution [(thf)<sub>4</sub>Li<sub>2</sub>(NtBu)<sub>4</sub>S] (201 mg, 0.296 mmol, 1.0 eq.) in THF (2 mL) was added and stirred overnight. After lithium chloride was filtered off and the brown solution was stored at –24 °C, colorless crystals were obtained after 4 days. Yield: 71 mg, 0.102 mmol, 34%; Elemental analysis (found (calc.) [%]): C 50.97 (51.96), H 9.34 (10.17), N 9.89 (10.10), S 4.76 (4.62). <sup>1</sup>H-NMR (300 MHz, THF-d<sub>8</sub>): δ (ppm) = 3.59–3.57 (m, 8 H, O(CH<sub>2</sub>)<sub>2</sub>), 1.74–1.71 (m, 8 H, O(CH<sub>2</sub>)<sub>2</sub>(CH<sub>2</sub>)<sub>2</sub>), 1.30 (s, 52 H, CH<sub>3</sub>); <sup>7</sup>Li-NMR (300 MHz, THF-d<sub>8</sub>): δ (ppm) = 0.18 (s, 1 Li); <sup>13</sup>C-NMR (300 MHz, THF-d<sub>8</sub>): δ (ppm) = 67.5 (s, 4 O(CH<sub>2</sub>)<sub>2</sub>), 57.6 (s, 4 CCH<sub>3</sub>), 30.4 (s, 18 CCH<sub>3</sub>), 26.3 (s, 4 O(CH<sub>2</sub>)<sub>2</sub>(CH<sub>2</sub>)<sub>2</sub>); <sup>29</sup>Si-NMR (300 MHz, THF-d<sub>8</sub>): δ = –113.0 (s, 2 Si).

### Single-crystal structural analysis

Single crystals were selected from a Schlenk flask under argon atmosphere and covered with perfluorated polyether oil on a microscope slide, which was cooled with a nitrogen gas flow supplied by the X-TEMP2 device.<sup>32</sup> An appropriate crystal was selected using a polarizing microscope, fixed on the tip of a MiTeGen<sup>®</sup> MicroMount, transferred to a goniometer head, and shock cooled by the crystal cooling device. The data for **3**, **4**, and **5** were collected from these shock-cooled crystals at 100(2) K. The data for **3** and **4** were collected on an Incoatec Mo microfocus source<sup>33</sup> equipped with Helios mirror optics and an APEX II detector at a D8 goniometer. The data for **5** were measured on a Bruker TXS Mo rotating anode with Helios mirror optics and an APEX II detector at a D8 goniometer. Important data are summarized in Table 1. Both diffractometers used Mo K<sub>α</sub> radiation, λ = 0.71073 Å. The data for all structures were integrated with SAINT,<sup>34</sup> and an empirical absorption correction (SADABS)<sup>35</sup> was applied. The structures were solved by direct methods (SHELXS-97)<sup>36</sup> and refined by full-matrix least-squares methods against *F*<sup>2</sup> (SHELXL-97)<sup>19,37</sup> within the SHELXLE GUI.<sup>38</sup> All non-hydrogen atoms were refined with anisotropic displacement parameters. The hydrogen



Table 1 Crystal and structure refinement parameters for compounds 3, 4, and 5

Parameters	3	4	5
CCDC no.	1011527	1011528	1011529
Empirical formula	C <sub>34</sub> H <sub>66</sub> Cu <sub>2</sub> N <sub>4</sub> O <sub>6</sub> S	C <sub>52</sub> H <sub>108</sub> Cd <sub>2</sub> I <sub>4</sub> Li <sub>2</sub> N <sub>4</sub> O <sub>9</sub> S	C <sub>30</sub> H <sub>70</sub> LiN <sub>5</sub> O <sub>2</sub> SSi <sub>2</sub> Zn
Formula weight	786.04	1711.76	693.46
Crystal system	Monoclinic	Monoclinic	Monoclinic
Space group	<i>P2<sub>1</sub>/n</i>	<i>P2<sub>1</sub>/c</i>	<i>C2/c</i>
<i>a</i> /Å	12.546(2)	15.428(2)	15.507(3)
<i>b</i> /Å	9.281(2)	22.442(3)	15.390(3)
<i>c</i> /Å	16.950(3)	20.869(2)	18.081(3)
$\beta$ /°	93.85(2)	103.56(2)	113.090(10)
<i>V</i> /Å <sup>3</sup> , <i>Z</i>	1969.2(6), 2	7024.2(16), 4	3969.4(13), 4
Density (calcd)	1.326 g cm <sup>-3</sup>	1.619 g cm <sup>-3</sup>	1.160 g cm <sup>-3</sup>
Absorption coefficient	1.178 mm <sup>-1</sup>	2.440 mm <sup>-1</sup>	0.762 mm <sup>-1</sup>
<i>F</i> (000)	840	3400	1512
Crystal size/mm	0.08 × 0.08 × 0.01	0.10 × 0.10 × 0.05	0.05 × 0.05 × 0.02
$\theta$ range for data collection	1.958 to 26.371°	1.353 to 25.521°	1.947 to 27.222°
Limiting indices	-15 ≤ <i>h</i> ≤ 15; -11 ≤ <i>k</i> ≤ 11; -21 ≤ <i>l</i> ≤ 21	-18 ≤ <i>h</i> ≤ 18; -27 ≤ <i>k</i> ≤ 27; -25 ≤ <i>l</i> ≤ 25	-19 ≤ <i>h</i> ≤ 19; -19 ≤ <i>k</i> ≤ 19; -23 ≤ <i>l</i> ≤ 23
Reflections collected	26 841	284 967	41 968
Independent reflections	4036 ( <i>R</i> <sub>int</sub> = 0.0350)	12 944 ( <i>R</i> <sub>int</sub> = 0.1384)	4420 ( <i>R</i> <sub>int</sub> = 0.0822)
Completeness to $\theta$	100% ( $\theta = 25.242^\circ$ )	100.0% ( $\theta = 25.242^\circ$ )	100% ( $\theta = 25.242^\circ$ )
Refinement method	Full-matrix least-squares on <i>F</i> <sup>2</sup>	Full-matrix least-squares on <i>F</i> <sup>2</sup>	Full-matrix least-squares on <i>F</i> <sup>2</sup>
Data/restraints/parameters	4036/558/311	12 944/2738/883	4420/0/201
Goodness-of-fit on <i>F</i> <sup>2</sup>	1.045	1.062	1.053
Final <i>R</i> indices [ <i>I</i> > 2σ( <i>I</i> )]	<i>R</i> <sub>1</sub> = 0.0270, <i>wR</i> <sub>2</sub> = 0.0641	<i>R</i> <sub>1</sub> = 0.0510, <i>wR</i> <sub>2</sub> = 0.1090	<i>R</i> <sub>1</sub> = 0.0308, <i>wR</i> <sub>2</sub> = 0.0742
<i>R</i> indices (all data)	<i>R</i> <sub>1</sub> = 0.0367, <i>wR</i> <sub>2</sub> = 0.0673	<i>R</i> <sub>1</sub> = 0.0724, <i>wR</i> <sub>2</sub> = 0.1226	<i>R</i> <sub>1</sub> = 0.0438, <i>wR</i> <sub>2</sub> = 0.0797
Largest diff. peak and hole/e Å <sup>-3</sup>	0.416 and -0.259	1.886 and -1.933	0.384 and -0.353

atoms were refined isotropically on calculated positions using a riding model with their *U*<sub>iso</sub> values constrained to equal 1.5 times the *U*<sub>eq</sub> of their pivot atoms for terminal sp<sup>3</sup> carbon atoms and 1.2 times for all other carbon atoms. Disordered moieties were refined using bond lengths and angles restraints and anisotropic displacement parameter restraints. Crystallographic data (excluding structure factors) for the structures reported in this paper have been deposited with the Cambridge Crystallographic Data Centre. The CCDC numbers, crystal data and experimental details for the X-ray measurements are listed in Table 1.

## Conclusion

The three metal complexes [(*acac*)<sub>2</sub>Cu<sub>2</sub>(*NtBu*)<sub>4</sub>S] (3), [Li(*thf*)<sub>4</sub>]<sub>2</sub>[I<sub>4</sub>Cd<sub>2</sub>(*NtBu*)<sub>4</sub>S] (4) and [(*thf*)<sub>2</sub>Li{(SiMe<sub>3</sub>)<sub>2</sub>N}Zn(*NtBu*)<sub>4</sub>S] (5) show that transition metal complexes containing the tetraimidosulfate dianion are feasible. With the right metal moiety at opposite sides of the tetrahedron they are stable and not subject to ligand scrambling. 4 can be envisaged as an intermediate on the metathesis reaction or a co-complex between CdI<sub>2</sub> and the lithium precursor.<sup>39</sup> Like in the intriguing structure of [(*thf*)<sub>2</sub>Li{(NtBu)<sub>3</sub>SMe}-ZnMe<sub>2</sub>]<sup>26</sup> the S–N bonds vary considerably in the heterobimetallic complex [(*thf*)<sub>2</sub>Li{(SiMe<sub>3</sub>)<sub>2</sub>N}-Zn(*NtBu*)<sub>4</sub>S] (5). The electropositive sulfur atom inside the imido nitrogen tetrahedron responds to the polarization induced by the coordinated metals. The more the N atoms lose density to the most electropositive metal at the outside the less they are attractive to the sulfur and the longer the S–N

bonds get. The electropositive sulfur in the inside mirrors the electron density distribution on the outside.

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## Notes and references

- H. W. Roesky, *Angew. Chem., Int. Ed. Engl.*, 1979, **18**, 91–97.
- T. Chivers, *Chem. Rev.*, 1985, **85**, 341–356.
- R. Mews, P. G. Watson and E. Lork, *Coord. Chem. Rev.*, 1997, **158**, 233–273.
- R. Fleischer and D. Stalke, *Coord. Chem. Rev.*, 1998, **176**, 431–450.
- D. Stalke, *Proc. Indian Acad. Sci.*, 2000, **112**, 155–170.
- J. K. Brask and T. Chivers, *Angew. Chem., Int. Ed.*, 2001, **40**, 3960–3976.
- F. T. Edelmann, S. Blaurock, V. Lorenz and T. Chivers, *Z. Anorg. Allg. Chem.*, 2008, **634**, 413–415.
- D. Stalke, *Chem. Commun.*, 2012, **48**, 9559–9573.
- M. M. Meinholz, E. Carl, E. Kriemen and D. Stalke, *Chem. Commun.*, 2011, **47**, 10948–10950.



- 10 M. M. Meinholz, S. K. Pandey, S. M. Deuerlein and D. Stalke, *Dalton Trans.*, 2011, **40**, 1662–1671.
- 11 M. M. Meinholz and D. Stalke, *Eur. J. Inorg. Chem.*, 2011, **2011**, 4578–4584.
- 12 M. M. Meinholz and D. Stalke, *Z. Anorg. Allg. Chem.*, 2011, **637**, 2233–2238.
- 13 M. M. Meinholz, E. Carl, E. Kriemen and D. Stalke, *Chem. Commun.*, 2011, **47**, 10948–10950.
- 14 M. M. Meinholz and D. Stalke, *Z. Naturforsch., B: Chem. Sci.*, 2011, **66**, 981.
- 15 R. Fleischer, S. Freitag and D. Stalke, *J. Chem. Soc., Dalton Trans.*, 1998, 193–198.
- 16 R. Fleischer, A. Rothenberger and D. Stalke, *Angew. Chem., Int. Ed. Engl.*, 1997, **36**, 1105–1107.
- 17 R. Fleischer, B. Walfort, A. Gbureck, P. Scholz, W. Kiefer and D. Stalke, *Chem. – Eur. J.*, 1998, **4**, 2266–2274.
- 18 T. Chivers and J. Konu, *Comments Inorg. Chem.*, 2009, **30**, 131.
- 19 P. Müller, R. Herbst-Irmer, A. L. Spek, T. R. Schneider and M. R. Sawaya, in *Crystal Structure Refinement – A Crystallographer's Guide to SHELXL*, ed. P. Müller, Oxford University Press, Oxford, England, 2006.
- 20 D. Leusser, J. Henn, N. Kocher, B. Engels and D. Stalke, *J. Am. Chem. Soc.*, 2004, **126**, 1781–1793.
- 21 J. Henn, D. Ilge, D. Leusser, D. Stalke and B. Engels, *J. Phys. Chem. A*, 2004, **108**, 9442–9452.
- 22 M. S. Schmökel, S. Cenedese, J. Overgaard, M. R. V. Jørgensen, Y.-S. Chen, C. Gatti, D. Stalke and B. B. Iversen, *Inorg. Chem.*, 2012, **51**, 8607–8616.
- 23 R. D. Shannon, *Acta Crystallogr., Sect. A: Cryst. Phys., Diffr., Theor. Gen. Cryst.*, 1976, **32**, 751–768.
- 24 K. Heinze and A. Reinhart, *Inorg. Chem.*, 2006, **45**, 2695–2703.
- 25 D. R. Lide, *CRC Handbook of Chemistry and Physics*, Taylor & Francis, Boca Raton, 2011–2012, vol. 92.
- 26 B. Walfort, A. P. Leedham, C. R. Russell and D. Stalke, *Inorg. Chem.*, 2001, **40**, 5668–5674.
- 27 (a) D. R. Armstrong, D. Barr, W. Clegg, R. E. Mulvey, D. Reed, R. Snaith and K. Wade, *J. Chem. Soc., Chem. Commun.*, 1986, 869; reviews: (b) K. Gregory, P. v. R. Schleyer and R. Snaith, *Adv. Inorg. Chem.*, 1991, **37**, 47; (c) R. E. Mulvey, *Chem. Soc. Rev.*, 1991, **20**, 167; (d) R. E. Mulvey, *Chem. Soc. Rev.*, 1998, **27**, 339.
- 28 *Cambridge Structural Database, v5.35 (November 2013)*, Cambridge Crystallographic Data Centre, Cambridge, 2013.
- 29 N. E. Brese and M. O'Keeffe, *Acta Crystallogr., Sect. B: Struct. Sci.*, 1991, **47**, 192–197.
- 30 (a) W. Schlenk and A. Thal, *Ber. Dtsch. Chem. Ges.*, 1913, **46**, 2840–2854; (b) W. Schlenk, J. Appenrodt, A. Michael and A. Thal, *Ber. Dtsch. Chem. Ges.*, 1914, **47**, 473–490.
- 31 [http://www.stalke.chemie.uni-goettingen.de/virtuelles\\_labor/de.html](http://www.stalke.chemie.uni-goettingen.de/virtuelles_labor/de.html).
- 32 (a) T. Kottke and D. Stalke, *J. Appl. Crystallogr.*, 1993, **26**, 615–619; (b) T. Kottke, R. J. Lagow and D. Stalke, *J. Appl. Crystallogr.*, 1996, **29**, 465–468; (c) D. Stalke, *Chem. Soc. Rev.*, 1998, **27**, 171–178.
- 33 T. Schulz, K. Meindl, D. Leusser, D. Stern, J. Graf, C. Michaelsen, M. Ruf, G. M. Sheldrick and D. Stalke, *J. Appl. Crystallogr.*, 2009, **42**, 885–891.
- 34 *SAINT v7.68A in Bruker APEX v2011.9*, Bruker AXS Inst. Inc., Madison, USA, 2008.
- 35 G. M. Sheldrick, *SADABS 2008/2*, Universität Göttingen, Germany, 2008.
- 36 G. M. Sheldrick, *Acta Crystallogr., Sect. A: Fundam. Crystallogr.*, 1990, **46**, 467–473.
- 37 G. M. Sheldrick, *Acta Crystallogr., Sect. A: Fundam. Crystallogr.*, 2008, **64**, 112–122.
- 38 C. B. Hübschle, G. M. Sheldrick and B. Dittrich, *J. Appl. Crystallogr.*, 2011, **44**, 1281–1284.
- 39 (a) E. Hevia, J. Z. Chua, P. Garzía-Álvarez, A. R. Kennedy and M. D. McCall, *Proc. Natl. Acad. Sci. U. S. A.*, 2010, **107**, 5294–5299; (b) D. R. Armstrong, W. Clegg, P. Garzía-Álvarez, A. R. Kennedy, M. D. McCall, L. Russo and E. Hevia, *Chem. – Eur. J.*, 2011, **17**, 8333–8341.

