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# A sulfur-templated Ni–Ni' coordination polymer that relies on a polarizable nickel nitrosyl hub†

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The templating properties of a diaza-nickel-*cis*-dithiolate towards triphenylphosphine gold(I), yielding a *transoid*  $[\text{Ni}(\text{N}_2\text{S}_2)\cdot 2\text{Au}(\text{PPh}_3)]$  complex (T. A. Pinder, S. K. Montalvo, A. M. Lunsford, C.-H. Hsieh, J. H. Reibenspies and M. Y. Darensbourg, *Dalton Trans.*, 2014, **43**, 138–144) suggested that a suitable analogue of  $d^{10}$ -Au(I), *i.e.*,  $\{\text{Ni}(\text{NO})\}^{10}$ , could generate a tetrahedral nickel node for a  $[\text{Ni}(\text{N}_2\text{S}_2)\cdot 2\text{Ni}(\text{NO})(\text{X})]_n$  coordination polymer. Monomeric precursors, derived from Feltham's  $[(\text{Ph}_3\text{P})_2\text{Ni}(\text{NO})(\text{Cl})]$  (R. D. Feltham, *Inorg. Chem.*, 1964, **3**, 116–119) produced the bidentate/sulfur-chelated  $[\text{Ni}(\text{N}_2\text{S}_2)\cdot \text{Ni}(\text{NO})(\text{X})]$  species with loss of  $\text{PPh}_3$ . Exchange of  $\text{Cl}^-$  by azide,  $\text{N}_3^-$ , in the  $\{\text{Ni}(\text{NO})\}^{10}$  synthon led to the balance of electrophilicity at  $\text{Ni}(\text{NO})$  and non-covalent (H-bonding and van der Waals) interactions that stabilized the extended chain of bridging sulfurs, in *transoid* connectivities, between a square planar  $\text{Ni}^{\text{II}}$  and a tetrahedral Ni, the latter within the electronic and spin-delocalized  $\{\text{Ni}(\text{NO})\}^{10}$  system. This study defines a new path that creates coordination polymers using metallodithiolates, the success of which, in this case, depends on the highly polarizable  $\{\text{Ni}(\text{NO})\}^{10}$  unit.

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## Introduction

Coordination polymers are broadly defined as extended structures based on metal ions linked into an infinite chain, sheet, or three-dimensional construction by bridging ligands.<sup>1–4</sup> Once formed, the balance of electronic and steric features/requirements that lead to the thermodynamic preference of observed connections in coordination polymers, may be, in many cases, readily rationalized. However, *a priori* design principles for a particular set of synthons, are not so easily predicted. Thus, advances in this huge field are typically made from knowledge of serendipitous prior success(es), based on certain building blocks and their fundamental properties.

A broad range of molecular compositions and structures are known to result from the ability of metallo-*cis*-dithiolates (Fig. 1) to capture multiple and numerous exogenous metals at the thiolate sulfurs.<sup>5</sup> Various synthetic approaches continue to be developed for entry into hetero bi- and polymetallic complexes, particularly extensive for square-planar,  $d^8$   $\text{Ni}(\text{II})\text{N}_2\text{S}_2$  derivatives, where the  $\text{N}_2\text{S}_2$  platform is contiguous nitrogen and sulfur linked by hydrocarbons. This arrangement places the two thiolates in a propitious position for bidentate ligation to exogeneous metals, including  $\text{Ni}\cdots\text{Ni}'$  arrangements in

biology.<sup>6</sup> In general, the bonding within  $[\text{M}(\eta^2\text{-}\mu\text{-SR})_2]\text{M}'$  may be compared to diphosphine- $\text{M}'$  or bipyridine- $\text{M}'$  interactions, as the nucleophilicity of metal-bound thiolate sulfurs has been somewhat deactivated relative to anionic  $\text{RS}^-$ , making the  $\text{M-S-R}$  similar to neutral P or N donors.<sup>5</sup> However, the stereochemistry demanded by the remaining sulfur lone pairs, and the orientation of their orbital lobes in MSR, particularly *cis*- $\text{M}(\text{SR})_2$ , provide structural possibilities (as shown in Fig. 1) unavailable to phosphorus or nitrogen donors. Most importantly, they generate stable sulfur-bridged polymetallics.

The successful linking of  $\text{NiN}_2\text{S}_2$  molecules through the attachment of electrophiles to the metal-bound thiolate sulfurs is an obvious strategy that offers an enormous prospect for synthesis of S-based coordination polymers. Most structural platforms involving  $[\text{M}(\eta^2\text{-}\mu\text{-SR})_2]\text{M}'$  thus far reported are related to bidentate  $\text{M}(\eta^2\text{-}\mu\text{-SR})_2$  binding in which the stereochemistry of the  $[\text{M}(\eta^2\text{-}\mu\text{-SR})_2]\text{M}'$  is governed by the minimization of sulfur lone pair repulsion.<sup>7</sup> The *cis*-configuration generates a butterfly-type hinge at the bridge (Fig. 1, EROPEG).<sup>8</sup> However, the placement of two  $\text{M}'$  electrophiles in a *transoid* arrangement relative to the  $\text{M}(\mu\text{-SR})_2$  plane, observed for a bis- $[\text{Au}(\text{PPh}_3)]_2$  derivative (Fig. 1, TINDUR) hints at the opportunity for designing coordination polymers based on  $\text{NiN}_2\text{S}_2$  bridges that link metal units in possession of two accessible open sites.<sup>9</sup>

It is well known that sulfur-based linkers exhibit energetic alignment with transition metal d orbitals, leading to stronger interactions and potentially superior material properties.<sup>10</sup> A prominent example from the Anderson group has revealed that a 4Fe4S-based coordination polymer can be used as a

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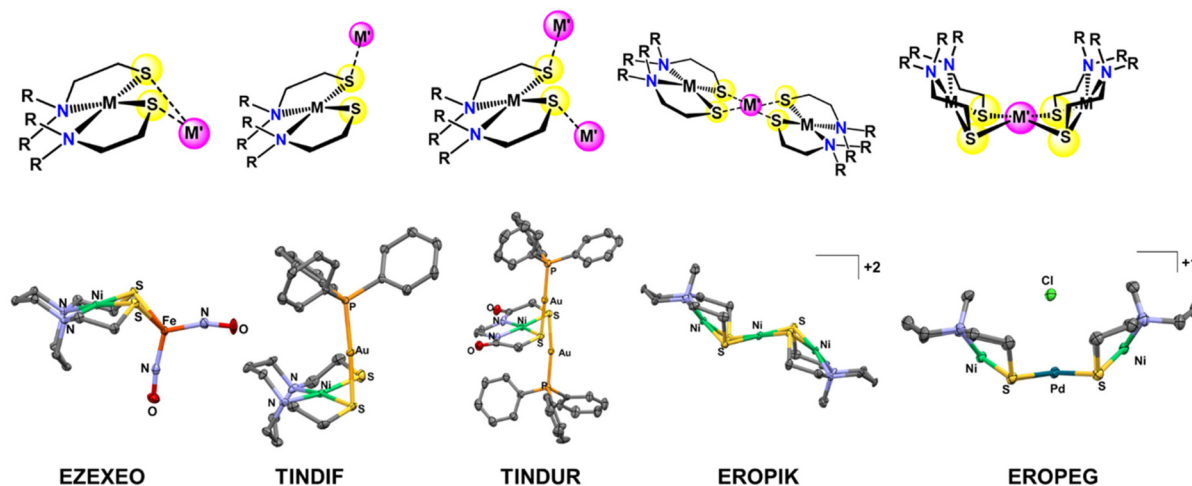


Fig. 1 Examples of various binding modes of  $\text{NiN}_2\text{S}_2$  due to presence of two lone pairs on each sulfur atom.<sup>8,9,14</sup> Identifiers taken from deposits in Cambridge Crystallographic Data Base.

heterogeneous, proton–electron coupled transfer mediator during electrocatalysis.<sup>11</sup> These highly active and robust cocatalysts can be easily recyclable during the electrocatalysis as compared to the monomeric  $4\text{Fe}_4\text{S}$  clusters.

Here we report a synthon,  $[\text{Ni}(\text{NO})(\text{Cl})(\text{PPh}_3)_2]$ ,<sup>12</sup> with  $\{\text{Ni}(\text{NO})\}^{10}$  as an electronic analogue of  $d^{10} \text{Au}^+$  or  $\{\text{Fe}(\text{NO})_2\}^{10}$  (*i.e.*, the total number of electrons in the metal-d orbitals +  $\pi^*$  manifold of NO is 10, represented in the Enemark–Feltham (E–F) notation<sup>13</sup>) which can bind with  $\text{NiN}_2\text{S}_2$  similar to the  $[\text{Ni}(\text{N}_2\text{S}_2)\cdot\text{Fe}(\text{NO})_2]$  complex, EZEXEO in Fig. 1.<sup>14</sup> Upon replacement of chloride by azide, the bidentate binding of the  $\text{NiN}_2\text{S}_2$  opens up and exposes an open site on the neutral  $\text{Ni}(\text{NO})(\text{N}_3)$  unit. This unit is found to serve as a hub for connecting the  $\text{NiN}_2\text{S}_2$  linkers in transoid arrangement as seen for the bis- $[\text{Au}(\text{PPh}_3)_2]$  derivative, TINDUR, Fig. 1.<sup>9</sup> The successful conversion of the azide derivative into the metallo polymer contrasts to the analogous  $[\text{Ni}(\text{NO})(\text{Cl})(\text{PPh}_3)_2]$  which is incapable of such development; the latter serves as a basis for understanding the balance of nickel electrophilicity and geometrical preferences. This report defines a new path to construct S-based coordination polymers using available lone pairs of sulfur in metallodithiolate ligands, the success of which is based on the electronically and structurally responsive  $\{\text{Ni}(\text{NO})\}^{10}$  unit.

## Results and discussion

### Synthesis and characterization

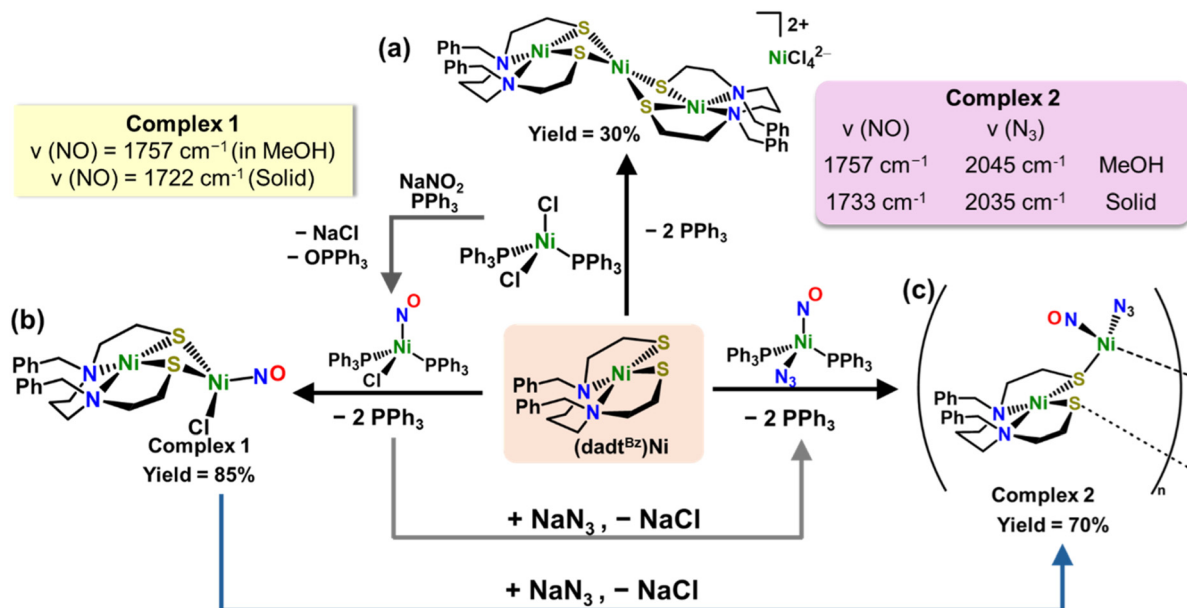
*(Caution! All described syntheses of azides result in potential energetic compounds with possible sensitivities towards heat, friction, impact and discharge. Although no incidents occurred during these syntheses personal safety equipment as well as grounded equipment is strongly recommended! Only small quantities (10–20 mg) of material should be used (at a*

time) during reactions with azide and metal spatulas should never be used.)

Relative to phosphines, bipyridines or isonitriles, the sulfur lone pairs in  $\text{MN}_2\text{S}_2$  metallodithiolates are of greater nucleophilicity and make them reasonable synthons in the design of sulfur-bridged heterometallics, some of which are directed towards catalysis.<sup>15</sup> Reaction of  $(\text{dadt}^{\text{Bz}})\text{Ni}$  ( $\text{dadt}^{\text{Bz}}$  is *N,N'*-dibenzyl-3,7-diazanonane-1,9-dithiolate)<sup>16</sup> with  $\text{NiCl}_2(\text{PPh}_3)_2$  did not however produce the expected  $[(\text{dadt}^{\text{Bz}})\text{Ni}\cdot\text{NiCl}_2]$  adduct. Instead, two  $\text{Cl}^-$  ions were displaced, giving the ubiquitous trinickel complex with three square planes arranged in the stair-step configuration,  $[(\text{dadt}^{\text{Bz}})\text{Ni}\cdot\text{Ni}^{\text{II}}\cdot\text{Ni}(\text{dadt}^{\text{Bz}})]^{2+}$ , having  $[\text{NiCl}_4]^{2-}$  as an anion, shown in Scheme 1a. The XRD-derived structure of this salt is shown in Fig. S1 and S2 and selected bond angles and lengths are given in Table S1.† Isolation of this complex prompted us to pursue the reported  $[\text{Ni}(\text{NO})(\text{Cl})(\text{PPh}_3)_2]$  complex as synthon.<sup>12</sup>

The tetrahedral  $[\text{Fe}(\text{CO})_2(\text{NO})_2]$  synthon binds with  $\text{MN}_2\text{S}_2$  with loss of carbonyl ligands as shown in EZEXEO, Fig. 1.<sup>14</sup> We hypothesized that nitric oxide (NO) might be a suitable candidate to accommodate the additional electron density produced from binding of the metallodithiolates while maintaining the geometry of the tetrahedral Ni center of  $[\text{Ni}(\text{NO})(\text{Cl})(\text{PPh}_3)_2]$ . Accordingly, a stoichiometric amount of the  $\text{Ni}(\text{NO})$  reagent was added to a methanolic solution of  $(\text{dadt}^{\text{Bz}})\text{Ni}$  producing a colour change from brown to dark blue, concomitant with the loss of the  $\nu(\text{NO})$  absorption of the precursor complex  $[\text{Ni}(\text{NO})(\text{Cl})(\text{PPh}_3)_2]$  in  $\text{CH}_3\text{OH}$  at  $1735 \text{ cm}^{-1}$ ; a new band appeared at *ca.*  $1757 \text{ cm}^{-1}$  (Fig. S3a†). The reaction solution was stirred for an additional hour. Evaporation of the solvent and recrystallization from  $\text{CH}_3\text{CN}/\text{Et}_2\text{O}$  yielded complex **1** in 85% yield as a dark green crystalline solid suitable for X-ray diffraction, Scheme 1b. The solid state  $\nu(\text{NO})$  IR spectrum is shown in Fig. S3b.† Mass spectral analysis of this complex shows  $m/z = 518.0360$  which confirms the presence of the expected neutral complex **1** (Fig. S4 and S5†). Another signal at





**Scheme 1** Synthetic routes to complexes reported in this study. MeOH was used as a solvent for all reactions. Note: complex 2 is the coordination polymer, alternatively referred to as CP2.

$m/z = 780.1271$  was observed, interpreted as due to the presence of a di-nickel complex with one triphenyl phosphine *i.e.*,  $[(\text{dadtbz})\text{Ni}\cdot\text{Ni}(\text{NO})(\text{Cl})(\text{PPh}_3)_2]$ , analogous to a previously reported gold complex (Fig. 1, identification code: TINDIF).<sup>9</sup> Despite several attempts this complex could not be isolated or characterized further. However, a change in the  $\text{N}_2\text{S}_2$  ligand backbone led to success.

A  $[(\text{N}_2\text{S}_2)\text{Ni}\cdot\text{Ni}(\text{NO})(\text{Cl})(\text{PPh}_3)_2]$  complex similar to the Ni-Au TINDIF complex<sup>9</sup> was isolated and crystallographically characterized upon changing the Ni synthon from  $(\text{dadtbz})\text{Ni}$  to  $(\text{bme-dame})\text{Ni}$  (*bme-dame* = *N,N'*-dimethyl-*N,N'*-bis(2-mercapto-ethyl)-ethylenediamine) during reaction with  $[\text{Ni}(\text{NO})(\text{Cl})(\text{PPh}_3)_2]$ . The XRD structure of this complex is given in the ESI (Fig. S9 and S10<sup>†</sup>) and the corresponding mass spectrum is in Fig. S11 and S12.<sup>†</sup>

In order to remove the metal-bound chloride as well as to see if the structural platform can be expanded from complex 1, reaction with a stoichiometric amount of  $\text{NaN}_3$  was performed in methanol. Consistent with Feltman's reports, the  $[\text{Ni}(\text{NO})(\text{Cl})(\text{PPh}_3)_2]$  reacts with  $\text{NaN}_3$  to form  $[\text{Ni}(\text{NO})(\text{N}_3)(\text{PPh}_3)_2]$ .<sup>17</sup> The FTIR spectrum of the reaction mixture shows absorption bands at *ca.* 2045 and 1757  $\text{cm}^{-1}$ , Fig. S13.<sup>†</sup> While no shift was observed in the  $\nu(\text{NO})$  IR band at 1757  $\text{cm}^{-1}$ , the appearance of the new absorbance at 2045  $\text{cm}^{-1}$  is taken as convincing evidence of the presence of a metal-bound azide ( $\text{N}_3^-$ ) in the complex. Removal of methanol and applying crystallization conditions,  $\text{CH}_3\text{CN}/\text{Et}_2\text{O}$ , yielded dark greenish-brown coloured needles of complex 2 in 70% yield, Scheme 1c. The solid state IR spectrum of this complex shows  $\nu(\text{NO})$  and  $\nu(\text{N}_3)$  at *ca.* 1733  $\text{cm}^{-1}$  and 2035  $\text{cm}^{-1}$  (Fig. S14<sup>†</sup>). Complex 2 can alternatively be obtained upon reaction of  $(\text{dadtbz})\text{Ni}$  with  $[\text{Ni}(\text{NO})(\text{N}_3)(\text{PPh}_3)_2]$  and it can be isolated as crystals from DMF/

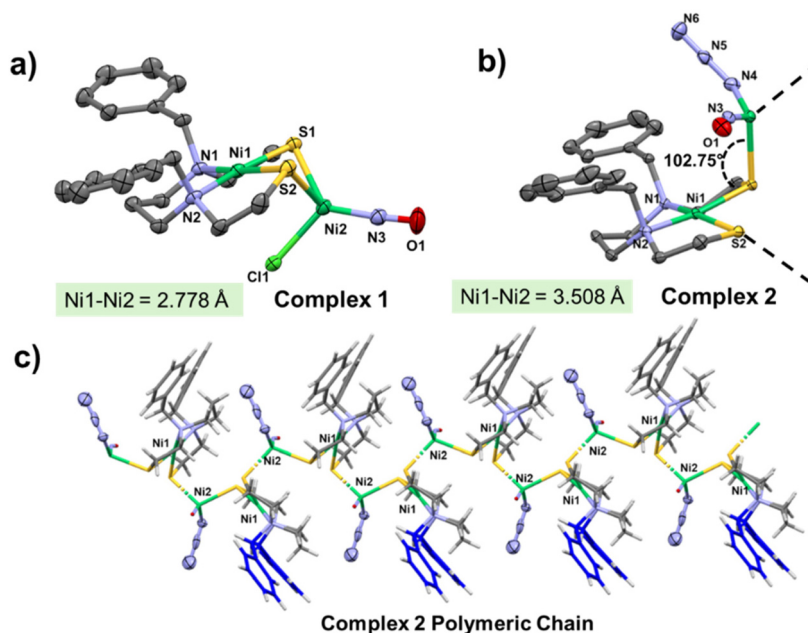
$\text{Et}_2\text{O}$  solution. X-ray diffraction analysis of the crystals obtained *via* both routes shows a one-dimensional coordination polymer chain. Detailed crystallographic description of this complex is given below. Once the complex crystallizes as a coordination polymer it becomes insoluble in almost all organic solvents (marginally in DMF). Note: we refer to this product as complex 2 throughout, although it is a coordination polymer.

### XRD characterization

The molecular structures of complexes 1,  $[(\text{dadtbz})\text{Ni}\cdot\text{Ni}(\text{NO})(\text{Cl})(\text{PPh}_3)_2]$ , and 2,  $[(\text{dadtbz})\text{Ni}\cdot\text{Ni}(\text{NO})(\text{N}_3)]_n$  are shown in Fig. 2 as 50% probability thermal ellipsoid plots; selected metric parameters are tabulated in Table S2.<sup>†</sup> Complexes 1 and 2 crystallize in the monoclinic  $P2_1/c$  and  $P2_1$  space groups respectively. The  $\text{Ni}_2\text{S}_2$  portion of complex 1, Ni1, is slightly distorted from square planar geometry with  $\tau_4 = 0.13$ . The second Ni center, Ni2, retains a distorted tetrahedral geometry with  $\tau_4 = 0.83$ . The NO molecule bound to Ni2 is almost linear,  $\angle\text{Ni2-N3-O1} = 175.40^\circ$ , *i.e.*, an increase in the  $\angle\text{Ni-N-O}$  bond angle from  $152.17^\circ$  (for monomeric  $[\text{Ni}(\text{NO})(\text{Cl})(\text{PPh}_3)_2]$ ) is observed upon displacement of the  $\text{PPh}_3$  groups by  $\text{NiN}_2\text{S}_2$ . A slight increase in the Ni-Cl bond length is also observed (from 2.27 Å in the Ni precursor to 2.31 Å in complex 1) upon coordination with the metallodithiolate ligand. The distance between the Ni centers in complex 1 is 2.778 Å. Based on the sum of the covalent atomic radii of Ni, this Ni-Ni' distance is beyond bonding. Crystal packing diagrams of this complex along three crystallographic axes are shown in Fig. S8,<sup>†</sup> and an expanded listing of pertinent metric data is provided in Table S2.<sup>†</sup>

Complex 2 crystallizes in the monoclinic  $P2_1$  space group. It can be crystallized from different solvents such as  $\text{CH}_3\text{CN}$ ,





**Fig. 2** Molecular structures of (a) complex 1  $[\text{Ni}(\text{dadt}^{\text{Bz}})\text{Ni}(\text{NO})(\text{Cl})]$  and (b) complex 2  $[\text{Ni}(\text{dadt}^{\text{Bz}})\text{Ni}(\text{NO})(\text{N}_3)]$  (asymmetric unit only is shown) showing 50% probability thermal ellipsoids. H-atoms are omitted for clarity. (c) The 1-D polymeric chain of 2.

MeOH as well as DMF. The following discussion refers to the crystals obtained from  $\text{CH}_3\text{CN}/\text{Et}_2\text{O}$ . X-ray crystallographic analysis of this complex reveals a one-dimensional coordination polymer structure containing repeating  $[[\text{dadt}^{\text{Bz}}\text{Ni}-\text{Ni}(\text{NO})(\text{N}_3)]]$  units, Fig. 2c. The unit cell of this species contains two asymmetric units, shown in Figs. S17 and S18.†

The nickel center inside the  $\text{N}_2\text{S}_2$  pocket in complex 2 is, as was found for complex 1, in a slightly distorted square planar geometry with  $\tau_4 = 0.18$ ; the displacement of the nickel from the mean  $\text{N}_2\text{S}_2$  plane (the plane created by  $\text{N}1\text{N}2\text{S}2$ ) is 0.015 Å. The other Ni center, Ni2 of the  $\{\text{Ni}(\text{NO})(\text{N}_3)\}$  unit, adopts a distorted tetrahedral geometry with  $\tau_4 = 0.84$ ; it is coordinated by S atoms of two distinct metallodithiolate ligands, as well as an NO and an azide,  $\text{N}_3^-$ .

The  $\angle\text{Ni}1-\text{S}1-\text{Ni}2$  angle in the CP 2 is  $102.75^\circ$ , in contrast to  $76.10^\circ$  in case of the dinickel complex 1. Upon coordination of the azide unit the metal nitrosyl ( $\angle\text{M}-\text{N}-\text{O}$ ) became significantly bent from  $175^\circ$  (seen for complex 1) to  $161^\circ$  ( $\angle\text{Ni}2-\text{N}3-\text{O}1 = 161.22^\circ$  for 2). A closer look on the structure of 2 suggests that the oxygen atom of NO molecule is within hydrogen bonding distance (2.5–2.6 Å) from the aliphatic and aromatic protons, Fig. 3d. The distance between the nickel centers increases substantially from 2.77 Å for complex 1, to 3.379 Å and 3.670 Å for the coordination polymer 2.

We examined the crystal packing of this complex for weak interactions that might assist in stabilizing this polymeric structure. As shown in Fig. 3a, the distance between the nickel centers in two adjacent layers is 12.175 Å (viewed along *c*-axis). The ‘zoomed-in’ view of the molecular structure shows that the azide group is oriented to indicate hydrogen-bonding interactions with phenyl hydrogens; four contacts are within distances of 2.7–3.1 Å, Fig. 3c.

There appears to be no stabilization by  $\pi-\pi$  stacking interactions between the aromatic rings<sup>18</sup> of the adjacent layers; instead, they are at an angle of *ca.*  $77^\circ$ , Fig. S19.† The arene planes in the same layer are separated by *ca.* 6.5 to 6.7 Å, Fig. S20.† In order to evaluate the weak interactions present in the coordination polymer, non-covalent interaction (NCI) analyses were performed on the unit cell of 2, see the Computational section below.

The solid-state structures of 2 obtained from different solvents ( $\text{CH}_3\text{CN}$ , MeOH and DMF) find almost identical metric parameters, except for slight differences in the Ni1–Ni2 distances and  $\angle\text{Ni}2-\text{N}3-\text{O}1$  angles, Fig. S22.† The Hirshfeld surface analysis of the crystals obtained from different solvents as well as that of complex 1 are given in the ESI (Fig. S37–40†).

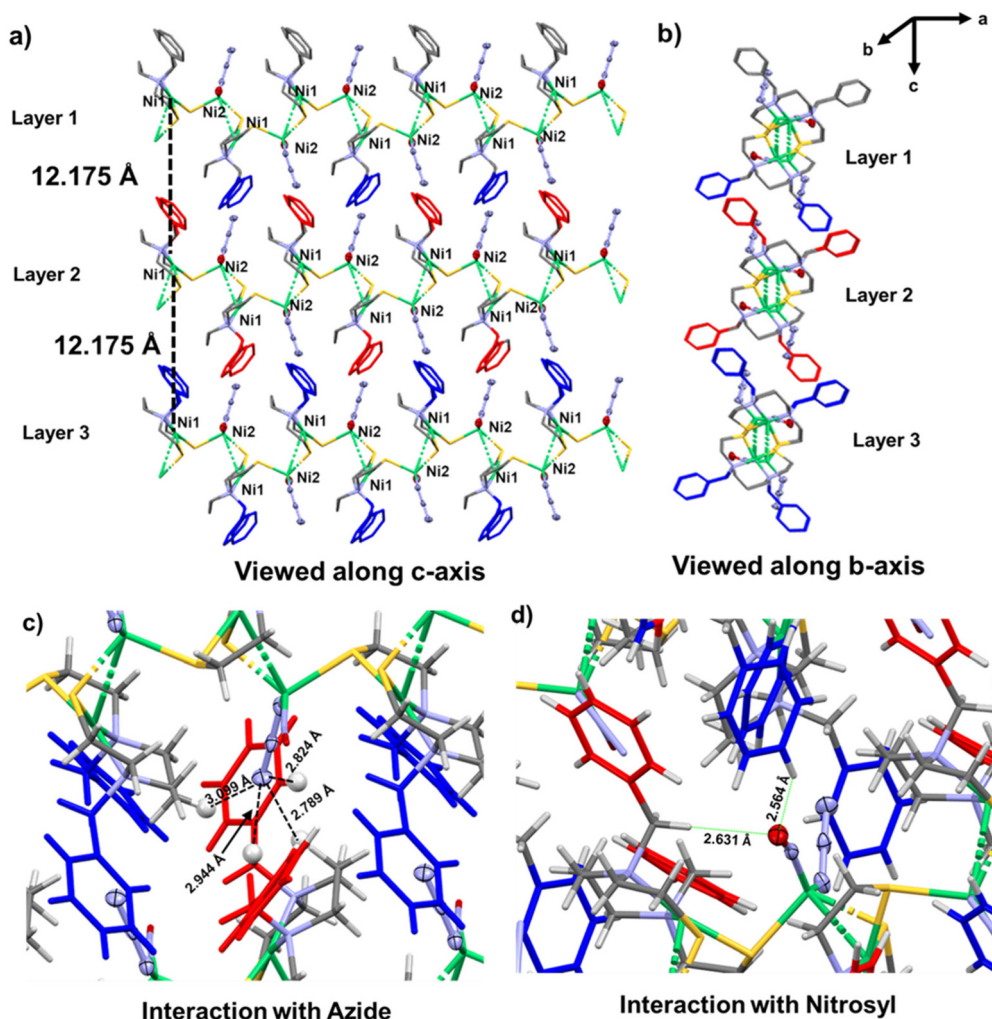
Crystallization from coordinating solvents like MeCN or DMF shows the presence of solvent molecules ( $\text{CH}_3\text{CN}/\text{DMF}$ ) inside the cavities created by stacking of multiple layers side by side, Fig. S22 and S26† (viewed along *b*-axis).

### Bulk purity and thermal stability

To reveal the phase purity of the coordination polymer (complex 2), a PXRD study was carried out on powdered samples within the  $2\theta$  range of  $5^\circ$ – $40^\circ$ , Fig. S31.† All significant peaks of the as-isolated powdered complex 2 were noted to be well-matched with the simulated pattern. The PXRD study along with elemental analysis of the coordination polymer confirms its bulk purity and crystallinity.

The coordination polymer is found to be stable under ambient conditions for weeks without any loss of NO or azide. Thermogravimetric analysis experiments on crystalline samples of 1 and 2 under  $\text{N}_2$  environment (Fig. S32†) indicated that complex 1 is stable up to  $150^\circ\text{C}$ , followed by complete





**Fig. 3** (a) Three layers of complex 2 viewed along *c*-axis. The distance between the Ni centers in two layers is 12.175 Å. (b) Three layers of complex 2 viewed along *b*-axis. (c) Zoomed in view of the interaction between the azide group and the aliphatic and aromatic hydrogens. (d) Zoomed in view of the interaction between nitrosyl group and aliphatic and aromatic hydrogens.

structural disruption beyond 150 °C. Complex 2 shows only 6.6 wt% decay up to *ca.* 141 °C, which can be interpreted as the loss of one mole MeCN per formula unit to yield the non-solvated 2. Complete structural disruption of the coordination polymer was observed after *ca.* 141 °C, similarly to complex 1.

### Redox properties

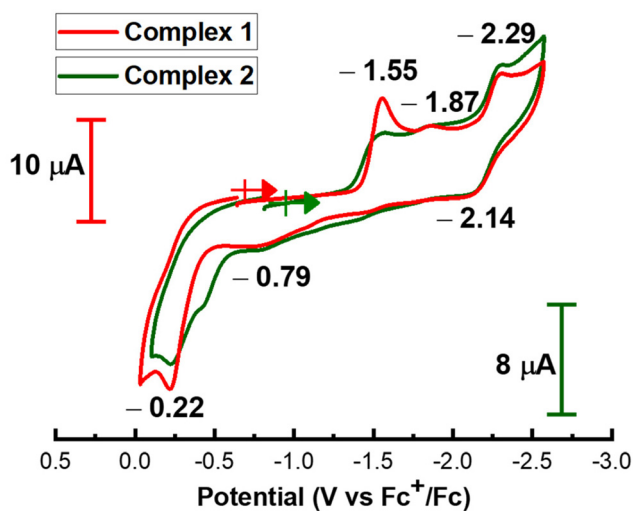
While the coordination polymer 2 is largely insoluble in CH<sub>3</sub>CN, its meager solubility in DMF permitted recording of the cyclic voltammograms of complexes, 1 and 2 at 22 °C under Ar atmosphere in DMF containing 0.1 M [*n*-Bu<sub>4</sub>N][PF<sub>6</sub>]<sub>4</sub> (Fig. S33<sup>†</sup>). Cyclic voltammetry of complex 1 was also attempted in MeCN, Fig. S35<sup>†</sup>. All values were referenced to internal Fc<sup>+</sup>/Fc at *E*<sub>1/2</sub> = 0.0 V and were recorded on sweeps initiated in the cathodic direction.

Three reduction events were observed in the initial cathodic sweep as shown, Fig. 4. The first irreversible reduction event at *ca.* -1.55 V for both complexes is assigned to the {Ni(NO)}<sup>10/</sup>

{Ni(NO)}<sup>11</sup> couple.<sup>19</sup> A second irreversible redox event observed at *ca.* -1.85 V in DMF is proposed to result from further reduction of the first reduced species. The third reduction event at *ca.* -2.30 V and its corresponding oxidation at -2.12 V is assigned to the reduction of the Ni<sup>2+</sup> center (Ni<sup>2+</sup> to Ni<sup>1+</sup>) housed within the N<sub>2</sub>S<sub>2</sub> ligand.

The lower solubility of the coordination polymer in DMF may explain the reduced current intensity in the CV features. Changes upon repeated scans of complex 2 are shown in Fig. S34<sup>†</sup>. Interestingly, both monomeric and polymeric complexes show almost identical redox behaviour in the cyclic voltammetry experiments which suggests the following: (a) the coordination polymer slightly dissociates in solution to give a monomeric complex similar to complex 1; or (b) the redox-activity of the coordination polymer is similar to that of the monomeric complex. The UV-vis spectra for the monomeric complex as well as the coordination polymer (sparingly soluble in DMF) (Fig. S36<sup>†</sup>) shows very similar absorption bands and





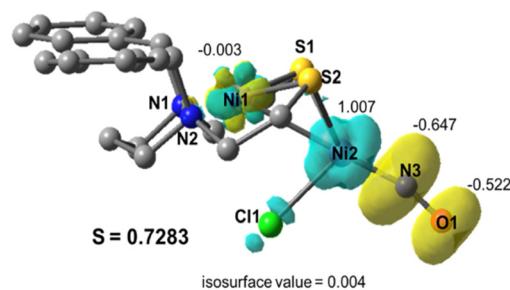
**Fig. 4** Cyclic voltammetry of complex 1 (red) and complex 2 (green) in DMF containing a 0.1 M [ $^t$ Bu<sub>4</sub>N][PF<sub>6</sub>] electrolyte and 2.0 mM complex at room temperature. Note that complex 2 has poor solubility in DMF, leading to reduced current intensity during the cyclic voltammetry analysis.

supports the cyclic voltametric result interpreted as the partial dissociation of the coordination polymer in DMF.

### Computational section

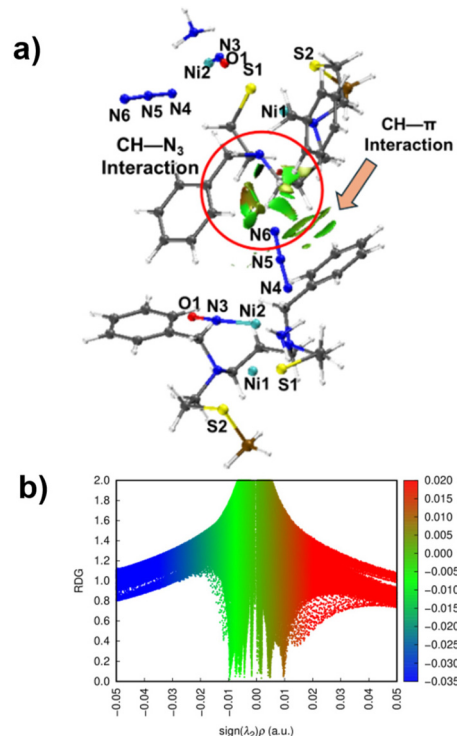
DFT Calculations were performed for the complex 1 with the TPSS<sup>20</sup> and B3LYP<sup>21</sup> functionals; the 6-31G(d,p) basis set<sup>22</sup> was used for C, H, N, O, S and Cl atoms whereas LANL2DZ basis set<sup>23</sup> was used for Ni atoms. The crystal structure of the complex was imported to use as the starting coordinates for the geometry optimization and subsequent vibrational frequency and energy calculations. All species were confirmed to be minimum energy structures by the absence of any imaginary frequencies. Experimentally, the known [Ni(NO)(L<sub>3</sub>)] complexes have been shown to have a diamagnetic ground state, which are best described as open-shell singlets (OSS) with a HS-Ni(II) center ( $S = 1$ ) antiferromagnetically coupled to a triplet state NO<sup>-</sup> ( $S = 1$ ) ligand.<sup>24</sup> Antiferromagnetic coupling for complex 1 was modelled in this manner with the B3LYP functional, and gives an overall spin of  $S_{\text{total}} = 0.73$ . Consistent with experiment, the <sup>1</sup>H NMR spectrum of this complex showed broad signals at room temperature (Fig. S6<sup>†</sup>). As expected, the single determinant wavefunction contains  $M_s = 0$  components from  $S > 0$  spin states. This optimized BS open-shell singlet state is found to be  $-7.59 \text{ kcal mol}^{-1}$  more stable than the corresponding closed-shell singlet structure. As shown by the wavefunction's spin density in Fig. 5, the majority of the spin is on Ni(II) and the NO<sup>-</sup> with spin populations of +1.01 and  $-1.16$ , respectively.

Interestingly, attempts to optimize the geometry of complex 1 as an open-shell singlet state with the TPSS functional converged to the closed-shell singlet state ( $S_{\text{total}} = 0$ ). The calculated structure for both methods are in accord with the experimental crystal structure shown in Fig. S40 and Table S4.<sup>†</sup>



**Fig. 5** Spin density plot for optimized BS-singlet structure of complex 1.

Parallel calculations for the monomeric form of azide, found as it exists in complex 1, as an OSS in B3LYP and as CSS in the TPSS (Table S4<sup>†</sup>), shows accumulation of electron density on NO which leads to decrease in the Ni–N–O bond angle. The  $\tau_4$  value for the Ni2 center decreases from the typical tetrahedral value toward the square planar value as shown in Table S4.<sup>†</sup> Apparently, these structural changes destabilize the monomeric form of the Ni(N<sub>2</sub>S<sub>2</sub>)-Ni(NO)(N<sub>3</sub>). Experimentally, the Ni–N–O bond is found to be more bent in the polymeric complex 2 than that of monomeric complex 1; nevertheless, the polymer has maintained the strong tetrahedral character of the complex 1 (see Table S2<sup>†</sup>).



**Fig. 6** (a) NCI index on the unit cell of complex 2 using iso surface value of 0.5 au and the color scale is  $-0.05 < \rho < 0.05$  au. CH– $\pi$  interactions are shown with arrow and the red circle indicates the CH–N<sub>3</sub> interaction. (b) Reduced density gradient plot shows prominent attractive interactions in complex 2.



Notably, the Ni–S bond distances are shorter in the polymeric complex than the monomeric complex. We assume this result may contribute to the stabilization of the polymer. In order to examine the non-covalent interactions (NCI) in the coordination polymer, **2**, the ground-state geometry of the assemblies found in the solid state was used to evaluate the attractive/repulsive character of the interactions by using Multiwfn to generate NCI Plots.<sup>25,26</sup> The Visual Molecular Dynamics (VMD) program was used to visualize the prominent interactions present in the molecule.<sup>27</sup> This methodology has been used for coordination polymers to estimate the contributions of a variety of interactions in the solid-state including hydrogen bonding,  $\pi$ – $\pi$  stacking and other weak van der Waals interactions.<sup>28</sup>

Since complex **2** is a coordination polymer, for investigations of limited views of units within the CP we terminated the S end with BH<sub>3</sub> and the Ni end with NH<sub>3</sub>. We performed a single point calculation on two asymmetric units of this “molecule” (two layers) which are adjacent to each other. The NCI plot of this construction indicates a prominent CH– $\pi$  interaction between the aromatic ring of one layer with the aromatic protons of the adjacent layer, Fig. 6. Hydrogen bonding and weak van der Waals interactions as indicated by the green iso-surface, are also observed between the azide group and the aliphatic and aromatic hydrogens (Fig. 6, red circle). The non-covalent interactions (NCI) coupled with strengthening of the Ni–S bonds help to stabilize the azide complex as a coordination polymer.

## Conclusion

This synthetic approach to a coordination polymer took advantage of the properties of two synthons:

(a) The templating ability of the *cis*-dithiolates in a Ni<sup>II</sup>(N<sub>2</sub>S<sub>2</sub>) complex that leads to the assembly of two strong electrophile/sulfur interactions transoid to each other, linking the Ni<sup>II</sup> with a second nickel imbedded within a polarizable and electroactive {Ni(NO)}<sup>10</sup> unit.

(b) The nitrosylated nickel source, with 10 electrons in its “golden” outer orbitals, is responsive to its ligand field. Its ability to electronically deform towards Ni<sup>II</sup>(NO)<sup>–</sup> is witnessed by shifts in crystallographic parameters. The application of Enemark–Feltham description for the overall electron count accounts for the oxidation state shifts within Ni(NO)<sup>10</sup>, *i.e.*, [Ni<sup>0</sup>(NO)<sup>+</sup>]  $\leftrightarrow$  [Ni<sup>I</sup>(NO)<sup>•</sup>]  $\leftrightarrow$  [Ni<sup>II</sup>(NO)<sup>–</sup>], and for the sensitive requirement of the bidentate binding ability of NiN<sub>2</sub>S<sub>2</sub> towards a single metal site, *vs.* its use as a bidentate bridging ligand that spans coordination sites of two metals.

This electronic deformation or polarization within the delocalized [Ni(NO)]<sup>+</sup> unit, *i.e.*, Ni(NO)<sup>10</sup>, accommodates the alternating assembly of Ni $\cdots$ Ni<sup>I</sup> in the coordination polymer. While the primary Ni–S–Ni<sup>I</sup> bonds in the polymer chains are strong, weak intermolecular interactions, identified in NCI plots, add to the stability of the polymer structure. The role of the substituents on the N<sub>2</sub>S<sub>2</sub> framework must be considered in this regard.

The potential of *cis*-dithiolates, particularly of nickel, for the assembly of heterobimetallic biomimetics and of catalysts is well developed.<sup>15</sup> However, to our knowledge this report is the first to explore and identify a route to coordination polymers based on this background. Given that libraries of MN<sub>2</sub>S<sub>2</sub> exist, there is potential for extensive development to expand the range of S-based coordination polymers and design of materials from their condensation.

## Author contributions

Manish Jana: experimental procedures, isolation and characterization (including theoretical calculations) of products, and design of project. M. J. Wrote the first and modified drafts. Michael B. Hall: design of computational approaches and overseer of results, interpretation and presentation. Marcetta Y. Darensbourg: design of project and interpretation/presentation of results. Contributed to writing of manuscript.

## Data availability

All underlying data are available in the article itself and in the ESI.†

## Conflicts of interest

There are no conflicts to declare.

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## References

- 1 K. Biradha, A. Ramanan and J. J. Vittal, *Cryst. Growth Des.*, 2009, **9**, 2969–2970.
- 2 W. L. Leong and J. J. Vittal, *Chem. Rev.*, 2011, **111**, 688–764.
- 3 C. S. Diercks, M. J. Kalmutzki, N. J. Diercks and O. M. Yaghi, *ACS Cent. Sci.*, 2018, **4**, 1457–1464.
- 4 L. S. Xie, G. Skorupskii and M. Dinca, *Chem. Rev.*, 2020, **120**, 8536–8580.
- 5 (a) J. A. Denny and M. Y. Darensbourg, *Chem. Rev.*, 2015, **115**, 5248–5273; (b) J. A. Denny and M. Y. Darensbourg, *Coord. Chem. Rev.*, 2016, **324**, 82–89.
- 6 (a) C. Darnault, A. Volbeda, E. J. Kim, P. Legrand, X. Vernède, P. A. Lindahl and J. C. Fontecilla-Camps, *Nat.*



- Struct. Mol. Biol.*, 2003, **10**, 271–279; (b) M. Can, F. A. Armstrong and S. W. Ragsdale, *Chem. Rev.*, 2014, **114**, 4149–4174; (c) R. E. Treviño and H. S. Shafaat, *Curr. Opin. Chem. Biol.*, 2022, **67**, 102110; (d) H. S. Shafaat, A. C. Manesis and A. Yerbulekova, *Acc. Chem. Res.*, 2023, **56**, 984–993.
- 7 P. Guerrero-Almaraz, M. Quiroz, D. R. Rodriguez, M. Jana, M. B. Hall and M. Y. Darensbourg, *ACS Org. Inorg. Au*, 2023, **3**, 393–402.
- 8 M. L. Golden, S. P. Jeffery, M. L. Miller, J. H. Reibenspies and M. Y. Darensbourg, *Eur. J. Inorg. Chem.*, 2004, **2004**, 231–236.
- 9 T. A. Pinder, S. K. Montalvo, A. M. Lunsford, C.-H. Hsieh, J. H. Reibenspies and M. Y. Darensbourg, *Dalton Trans.*, 2014, **43**, 138–144.
- 10 (a) R. Dong, P. Han, H. Arora, M. Ballabio, M. Karakus, Z. Zhang, C. Shekhar, P. Adler, P. S. Petkov, A. Erbe, S. C. B. Mannsfeld, C. Felser, T. Heine, M. Bonn, X. Feng and E. Cánovas, *Nat. Mater.*, 2018, **17**, 1027–1032; (b) A. J. Clough, N. M. Orchanian, J. M. Skelton, A. J. Neer, S. A. Howard, C. A. Downes, L. F. J. Piper, A. Walsh, B. C. Melot and S. C. Marinescu, *J. Am. Chem. Soc.*, 2019, **141**, 16323–16330; (c) T. Kusamoto and H. Nishihara, *Coord. Chem. Rev.*, 2019, **380**, 419–439; (d) D. Sheberla, L. Sun, M. A. Blood-Forsythe, S. Er, C. R. Wade, C. K. Brozek, A. Aspuru-Guzik and M. Dincă, *J. Am. Chem. Soc.*, 2014, **136**, 8859–8862; (e) M. G. Campbell, D. Sheberla, S. F. Liu, T. M. Swager and M. Dincă, *Angew. Chem., Int. Ed.*, 2015, **54**, 4349–4352; (f) L. Sun, C. H. Hendon, M. A. Minier, A. Walsh and M. Dincă, *J. Am. Chem. Soc.*, 2015, **137**, 6164–6167; (g) J. A. DeGayner, I.-R. Jeon, L. Sun, M. Dincă and T. D. Harris, *J. Am. Chem. Soc.*, 2017, **139**, 4175–4184; (h) M. Fourmigué, C. E. Uzelmeier, K. Boubekour, S. L. Bartley and K. R. Dunbar, *J. Organomet. Chem.*, 1997, **529**, 343–350; (i) L. Sun, T. Miyakai, S. Seki and M. Dincă, *J. Am. Chem. Soc.*, 2013, **135**, 8185–8188; (j) G. Givaja, P. Amo-Ochoa, C. J. Gómez-García and F. Zamora, *Chem. Soc. Rev.*, 2011, **41**, 115–147; (k) J. Xie, J.-N. Boyn, A. S. Filatov, A. J. McNeece, D. A. Mazziotti and J. S. Anderson, *Chem. Sci.*, 2020, **11**, 1066–1078; (l) L. Sun, M. G. Campbell and M. Dincă, *Angew. Chem., Int. Ed.*, 2016, **55**, 3566–3579.
- 11 (a) N. E. Horwitz, J. Xie, A. S. Filatov, R. J. Papoular, W. E. Shepard, D. Z. Zee, M. P. Grahn, C. Gilder and J. S. Anderson, *J. Am. Chem. Soc.*, 2019, **141**, 3940–3951; (b) N. Jiang, A. Darù, Š. Kunstelj, J. G. Vitillo, M. E. Czaikowski, A. S. Filatov, A. Wuttig, L. Gagliardi and J. S. Anderson, *J. Am. Chem. Soc.*, 2024, **146**, 12243–12252.
- 12 R. D. Feltham, *Inorg. Chem.*, 1964, **3**, 116–119.
- 13 J. H. Enemark and R. D. Feltham, *Coord. Chem. Rev.*, 1974, **13**, 339–406.
- 14 P. Ghosh, S. Ding, R. B. Chupik, M. Quiroz, C.-H. Hsieh, N. Bhuvanesh, M. B. Hall and M. Y. Darensbourg, *Chem. Sci.*, 2017, **8**, 8291–8300.
- 15 (a) C.-H. Hsieh, S. Ding, Ö. F. Erdem, D. J. Crouthers, T. Liu, C. C. L. McCrory, W. Lubitz, C. V. Popescu, J. H. Reibenspies, M. B. Hall and M. Y. Darensbourg, *Nat. Commun.*, 2014, **5**, 3684; (b) S. Ding, P. Ghosh, A. M. Lunsford, N. Wang, N. Bhuvanesh, M. B. Hall and M. Y. Darensbourg, *J. Am. Chem. Soc.*, 2016, **138**, 12920–12927.
- 16 M. Jana, X. Zheng, T. Le, M. Quiroz, P. Guerrero-Almaraz, D. J. Darensbourg and M. Y. Darensbourg, *Adv. Sci.*, 2024, **11**, 2307113.
- 17 J. H. Enemark, *Inorg. Chem.*, 1971, **10**, 1952–1957.
- 18 D. B. Ninković, J. P. Blagojević Filipović, M. B. Hall, E. N. Brothers and S. D. Zarić, *ACS Cent. Sci.*, 2020, **6**, 420–425.
- 19 S. Kundu, P. N. Phu, P. Ghosh, S. A. Kozimor, J. A. Bertke, S. C. E. Stieber and T. H. Warren, *J. Am. Chem. Soc.*, 2019, **141**, 1415–1419.
- 20 J. Tao, J. P. Perdew, V. N. Staroverov and G. E. Scuseria, *Phys. Rev. Lett.*, 2003, **91**, 146401.
- 21 A. D. Becke, *J. Chem. Phys.*, 1993, **98**, 1372–1377.
- 22 W. J. Hehre, R. Ditchfield and J. A. Pople, *J. Chem. Phys.*, 1972, **56**, 2257–2261.
- 23 (a) P. J. Hay and W. R. Wadt, *J. Chem. Phys.*, 1985, **82**, 299–310; (b) Y. Yang, M. N. Weaver and K. M. Merz Jr., *J. Phys. Chem. A*, 2009, **113**, 9843–9851.
- 24 K. Fujisawa, T. Kataoka, K. Terashima, H. Kurihara, F. de Santis Gonçalves and N. Lehnert, *Molecules*, 2023, **28**, 6206.
- 25 T. Lu and F. Chen, *J. Comput. Chem.*, 2012, **33**, 580–592.
- 26 E. R. Johnson, S. Keinan, P. Mori-Sánchez, J. Contreras-García, A. J. Cohen and W. Yang, *J. Am. Chem. Soc.*, 2010, **132**, 6498–6506.
- 27 W. Humphrey, A. Dalke and K. Schulten, *J. Mol. Graphics*, 1996, **14**, 33–38.
- 28 (a) S. K. Seth, *Crystals*, 2018, **8**, 455; (b) A. Gogoi, A. Das, A. Frontera, A. K. Verma and M. K. Bhattacharyya, *Inorg. Chim. Acta*, 2019, **493**, 1–13; N. A. Ashashi, M. Kumar, R. M. Gomila, A. Frontera, H. N. Sheikh and S. C. Sahoo, *J. Mol. Struct.*, 2022, **1254**, 132329; (c) N. A. Ashashi, M. Kumar, R. M. Gomila, A. Frontera, H. N. Sheikh and S. C. Sahoo, *J. Mol. Struct.*, 2022, **1254**, 132329; (d) S. Dutta, S. Chakraborty, M. G. B. Drew, A. Frontera and A. Ghosh, *Cryst. Growth Des.*, 2019, **19**, 5819–5828.

