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A combined experimental and computational study on supramolecular assemblies in hetero-tetranuclear nickel(II)–cadmium(II) complexes with N₂O₄-donor compartmental Schiff bases†

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Two new hetero-tetranuclear nickel(II)/cadmium(II) complexes, a cubane [(CH₃CO₂)₂Ni₂(L¹)₂Cd₂(NCS)₂] (**1**) and a linear tetramer [(DMSO)₂NiL²Cd(NCS)(μ_{1,3}-SCN)Ni(DMSO)L²Cd(NCS)₂] (**2**) (where H₂L¹ = *N,N'*-bis(3-methoxysalicylidene)propane-1,3-diamine and H₂L² = *N,N'*-bis(3-ethoxysalicylidene)propane-1,3-diamine are potential octadentate compartmental Schiff bases), were synthesized and characterized. The structures of both complexes were confirmed by single crystal X-ray diffraction studies. Complex **1** contained a Ni₂Cd₂O₄ cubane core, whereas complex **2** featured an end-to-end thiocyanate-bridged tetranuclear moiety. Furthermore, complex **1** showed C–H...H–C interactions, whereas a unique S...π interaction was observed in complex **2**. Theoretical studies were performed using several computational tools such as NBO and AIM analyses. Both complexes showed photoluminescence in DMSO medium at room temperature upon irradiation with ultraviolet light. The lifetimes of the excited states were ~27 ns.

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Introduction

The condensation of various diamines and salicylaldehyde gives H₂salen-type Schiff bases, which are very well-known chelating ligands to synthetic inorganic chemists for the preparation of di- and polynuclear complexes.¹ The di-negative anionic forms of these Schiff bases generally act as tetradentate N₂O₂-donor chelating ligands to produce mononuclear complexes,² which can cleverly be utilized as ligands (metallo-ligands) to coordinate a second metal ion by exploiting the interesting bridging ability of the phenoxo oxygen atoms, thereby forming multinuclear complexes.³ The use of 3-alkoxy-salicylaldehyde instead of salicylaldehyde itself is more beneficial as it produces compartmental octadentate Schiff bases (with inner N₂O₂ and outer O₄ cores), which can again be used efficiently to prepare various homo-polynuclear complexes.⁴ These high-nuclearity transition and non-transition metal complexes have attracted much attention for their unusual physical properties⁵ and relevance to biological function in

metalloproteins.⁶ Among these, phenoxo-bridged tetrameric nickel(II) clusters have been extensively studied to explore the relationship between their structural features and the strength of the magnetic exchange interaction between the nickel(II) clusters.⁷ Reports on the formation of heteronuclear tetramers, however, are relatively fewer in the literature.⁸

Supramolecular interactions have been shown to play an important role in structural biology and supramolecular chemistry.⁹ The harmonization of various biological and physico-chemical methods is frequently implemented by the amalgamation of many non-covalent supramolecular interactions.¹⁰ A good number of familiar, strong, directional non-covalent interactions, such as hydrogen bonding and halogen bonding, have been used to manage multi-component supramolecular assemblies.¹¹ In addition, π...π stacking, cation...π, C–H...π, lone pair...π and anion...π interactions are also extremely significant in this field¹² and can manage the structures of bio-molecules such as proteins and DNA, several host-guest systems, enzyme-substrate binding.¹³ They can also participate in crystal engineering, drug-receptor interactions, enzyme inhibition, protein folding, *etc.*¹⁴ As a matter of fact, S...π interactions are crucial in the mechanism of sulphide:quinone oxidoreductase.^{14d} Their omnipresent role in such diverse fields has attracted the attention of theoretical chemists to investigate and try to understand the nature of these weak non-covalent interactions, and, as a result, the importance of these no-covalent interactions have been analyzed by various

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researchers using combined theoretical and experimental methods.¹⁵

In the present work, two similar compartmental N_2O_4 -donor Schiff bases were used to prepare two hetero-tetranuclear nickel(II)/cadmium(II) complexes: a cubane and a linear tetramer with $Ni_2Cd_2O_4$ cores. These are the first examples of any hetero-tetranuclear nickel(II)/cadmium(II) complexes with H_2salen -type Schiff bases. The unconventional supramolecular interactions found in the crystal packing of these complexes were theoretically analyzed focusing attention on the remarkable $S \cdots \pi$ interactions and also on the $C-H \cdots H-C$ interactions. Herein, we would like to report the synthesis, characterization, X-ray crystal structure, photoluminescence properties and results of the DFT study on the supramolecular architectures of these two new hetero-tetranuclear nickel(II)/cadmium(II) complexes.

Experimental section

Nickel(II) thiocyanate tetrahydrate was prepared in the laboratory following the literature method.¹⁶ All the other materials were commercially available, reagent grade and used as purchased from Sigma-Aldrich without further purification.

Preparations

Preparation of the ligands

Preparation of H_2L^1 [N,N' -bis(3-methoxysalicylidene)propane-1,3-diamine]. A methanol solution (10 ml) of 3-methoxysalicylaldehyde (304 mg, 2 mmol) and 1,3-diaminopropane (0.13 ml, 1 mmol) was refluxed for *ca.* 1 h to form H_2L^1 . The ligand was not isolated and was used directly for the synthesis of complex 1.

Preparation of H_2L^2 [N,N' -bis(3-ethoxysalicylidene)propane-1,3-diamine]. This ligand was prepared using a similar method as that used for H_2L^1 except that 3-ethoxysalicylaldehyde (332 mg, 2 mmol) was used instead of 3-methoxysalicylaldehyde. It was also not isolated and was used directly for the synthesis of complex 2.

Preparation of the complexes

Preparation of $[(CH_3CO_2)_2Ni_2(L^1)_2Cd_2(NCS)_2]$ (1). A methanol (10 ml) solution of cadmium(II) acetate dihydrate (270 mg, 1 mmol) was added to the methanol solution (20 ml) of H_2L^1 and the resulting solution was stirred for 15 min. A methanol (10 ml) solution of nickel(II) thiocyanate tetrahydrate (250 mg, 1 mmol) was then added to it. The stirring was continued for about 2 h. Single crystals, suitable for X-ray diffraction, were obtained after 3–4 days on slow evaporation of the solution in open atmosphere.

Yield: 460 mg (72%); based on nickel(II). Anal. Calc. for $C_{44}H_{46}Cd_2Ni_2O_{12}S_2N_6$ (FW = 1257.21): C, 42.04; H, 3.69; N, 6.68; Found: C, 41.9; H, 3.5; N, 6.7%. FT-IR (KBr, cm^{-1}): 1630 (C=N); 2067 (C≡N). UV-VIS [$\lambda_{max}(nm)$] [$\epsilon_{max}(L mol^{-1} cm^{-1})$] (DMSO): 239 (2.55×10^4); 375 (3.4×10^3); 668 (5.86); 901 (13.84).

Preparation of $[(DMSO)_2NiL^2Cd(NCS)(\mu_{1,3}-SCN)Ni(DMSO)L^2Cd(NCS)_2]$ (2). This ligand was prepared using a similar method as that used for complex 1 except that H_2L^2 was used instead of H_2L^1 . A few drops of DMSO were added and the resulting solution was kept for crystallization. Single crystals, suitable for X-ray diffraction, were obtained after 3–4 days on slow evaporation of the solution in open atmosphere.

Yield: 570 mg (74%); based on nickel(II). Anal. Calc. for $C_{52}H_{66}Cd_2Ni_2O_{11}S_7N_8$ (FW = 1545.82): C, 40.40; H, 4.30; N, 7.25; Found: C, 40.2; H, 4.2; N, 7.3%. FT-IR (KBr, cm^{-1}): 1628 (C=N); 2034, 2068 (C≡N). UV-VIS [$\lambda_{max}(nm)$] [$\epsilon_{max}(L mol^{-1} cm^{-1})$] (DMSO): 236 (2.75×10^4); 378 (3.6×10^3); 672 (7.46); 902 (8.64).

Physical measurements

Elemental analyses (carbon, hydrogen and nitrogen) were performed using a Perkin Elmer 240C elemental analyzer. IR spectra in KBr (4500–500 cm^{-1}) were recorded with a Perkin Elmer Spectrum Two spectrophotometer. Electronic spectra in DMSO were recorded on a Perkin Elmer Lambda 35 UV-visible spectrophotometer. Steady-state photoluminescence spectra in DMSO were obtained with a Shimadzu RF-5301PC spectrofluorometer at room temperature. Time-dependent photoluminescence spectra were recorded using a Hamamatsu MCP photomultiplier (R3809) and were analyzed using IBHDAS6 software. Powder X-ray diffraction was performed on a Bruker D8 instrument with Cu K_α radiation. In this process, the complexes were ground with a mortar and pestle to prepare fine powders. The powders were then dispersed with alcohol onto a zero background holder (ZBH). The alcohol was allowed to evaporate to provide a nice, even coating of powder adhered to the sample holder.

X-ray crystallography

Suitable single crystals of complexes 1 and 2 were used for the data collection using a Bruker SMART APEX II diffractometer equipped with graphite-monochromated Mo K_α radiation ($\lambda = 0.71073 \text{ \AA}$) at 100 K. The molecular structures were solved by a direct method and refined by full-matrix least squares on F^2 using the SHELX-2014 package.¹⁷ Non-hydrogen atoms were refined using anisotropic thermal parameters. The hydrogen atoms were placed in their geometrically idealized positions and constrained to ride on their parent atoms. Multi-scan empirical absorption corrections were applied to the data using the program SADABS.¹⁸ Details of the crystallographic data and refinement details are given in Table 1.

The crystal structure of complex 1 was somewhat problematic. The unit cell included some highly disordered solvent molecules, which could not be modelled as discrete atomic sites. We employed PLATON/SQUEEZE to calculate the diffraction contribution of the solvent molecules and thereby were able to produce a set of solvent-free diffraction intensities. Details about the SQUEEZE procedure are given in the CIF file.



Table 1 Crystal data and refinement details of complexes **1** and **2**

Complex	1	2
Formula	C ₄₄ H ₄₆ Cd ₂ Ni ₂ O ₁₂ S ₂ N ₆	C ₅₂ H ₆₆ Cd ₂ Ni ₂ O ₁₁ S ₇ N ₈
Formula weight	1257.21	1545.82
Temperature (K)	100	100
Crystal system	Triclinic	Monoclinic
Space group	<i>P</i> $\bar{1}$	<i>P</i> 2 ₁ / <i>n</i>
<i>a</i> (Å)	15.313(10)	12.070(3)
<i>b</i> (Å)	15.334(1)	20.252(5)
<i>c</i> (Å)	23.219(16)	26.697(7)
α (°)	74.682(4)	90
β (°)	77.101(4)	93.663(1)
γ (°)	89.888(4)	90
<i>Z</i>	4	4
<i>d</i> _{calc} (g cm ⁻³)	1.632	1.577
μ (mm ⁻¹)	1.689	1.497
<i>F</i> (000)	2528	3144
Total reflections	70 609	93 652
Unique reflections	18 091	11 581
Observed data	14 053	8813
[<i>I</i> > 2 σ (<i>I</i>)]		
No. of parameters	1225	743
<i>R</i> (int)	0.067	0.049
<i>R</i> ₁ , <i>wR</i> ₂ (all data)	0.0944, 0.2010	0.0619, 0.1054
<i>R</i> ₁ , <i>wR</i> ₂ [<i>I</i> > 2 σ (<i>I</i>)]	0.0783, 0.1941	0.0416, 0.0942

Theoretical methods

The calculations of the non-covalent interactions were carried out using TURBOMOLE version 7.0¹⁹ using the M06-2X/def2-TZVP level of theory. To evaluate the interactions in the solid state, we used the crystallographic coordinates and optimized the position of the H atoms. In the large assemblies computed for complex **1**, the SVP basis set was used instead of the time-

consuming def2-TZVP basis set. This procedure and level of theory have been successfully used before to evaluate similar interactions.²⁰ The interaction energies were computed by calculating the difference between the energies of isolated monomers and their assembly. The interaction energies were corrected for the Basis Set Superposition Error (BSSE) using the counterpoise method.²¹

The Natural Bond Orbital (NBO) method²² was employed to analyze the charge-transfer interactions between the occupied and empty orbitals using the NBO-3.1 program included within the Gaussian-09 program. Bader's "Atoms in molecules" theory was used to study the interactions discussed herein by means of the AIMall calculation package.²³

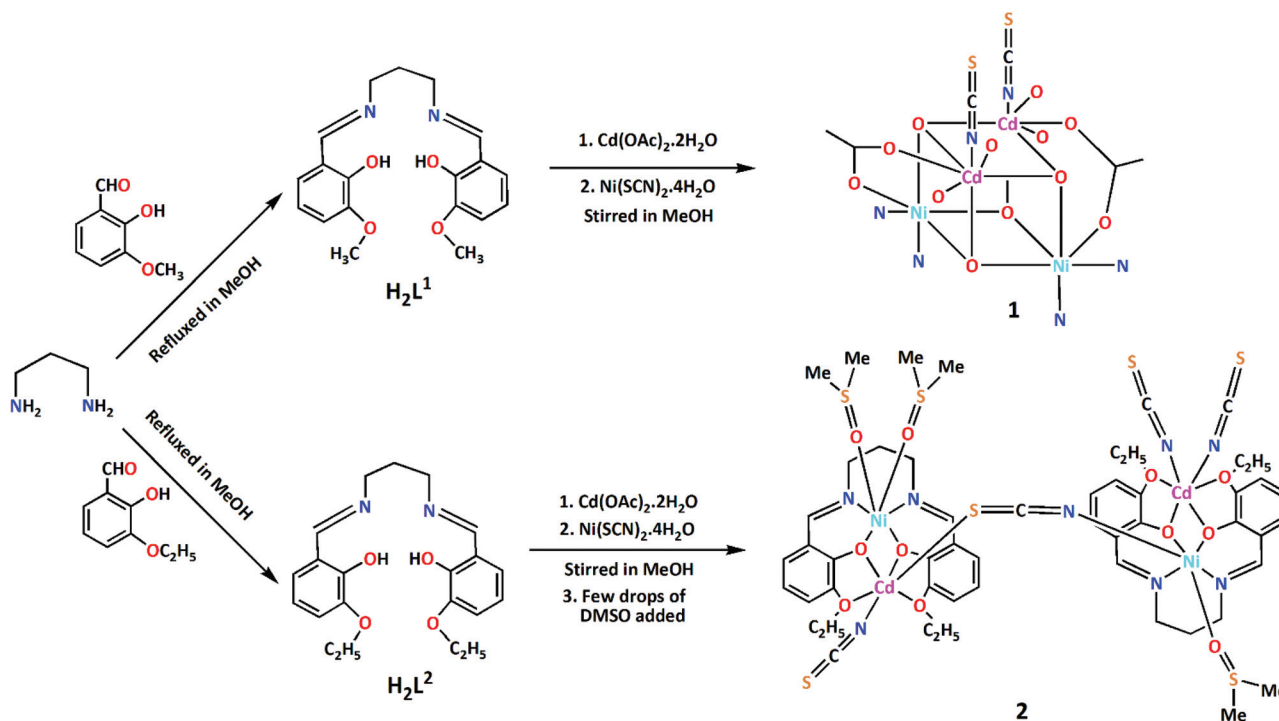
Results and discussion

Synthesis

1,3-Diaminopropane was refluxed with 3-methoxysalicylaldehyde and 3-ethoxysalicylaldehyde, respectively, to form two potential octadentate Schiff base ligands, H₂L¹ and H₂L², respectively, following the literature method.²⁴ These Schiff bases (H₂L¹ and H₂L²) on reaction with cadmium(II) acetate dihydrate, followed by the addition of nickel(II) thiocyanate tetrahydrate, gave two heteronuclear complexes. The formation of both complexes is shown in Scheme 1.

Description of the structures

[(CH₃CO₂)₂Ni₂(L¹)₂Cd₂(NCS)₂] (**1**). The X-ray crystal structure determination revealed that complex **1** crystallizes in the



Scheme 1 Preparation of the ligands and complexes. Only the cubane core of complex **1** is shown for clarity.



triclinic space group $P\bar{1}$. The complex contains two independent hetero-tetranuclear units (**A** and **B**) with equivalent geometries. A perspective view of unit **A** is shown in Fig. 1. The cubane core and the coordinating atoms are highlighted in Fig. 2. The cubane core of unit **B** has a similar structure, as shown in Fig. S1 {ESI†}. The important bond lengths of unit **A** are given in Table 2. The important bond lengths of unit **B** and the bond angles (units **A** and **B**) are gathered in Tables S1–S3.†

The molecular structure of unit **A** is built from isolated heteronuclear molecules of $[(\text{CH}_3\text{CO}_2)_2\text{Ni}_2(\text{L}^1)_2\text{Cd}_2(\text{NCS})_2]$, in which both nickel(II) centres are hexacoordinated having pseudo-octahedral geometries while cadmium(II) centres are heptacoordinated. H_2L^1 is a potential octadentate compartmental Schiff base with inner N_2O_2 and outer O_4 compartments. Nickel(II) centres occupy the inner N_2O_2 compartment,

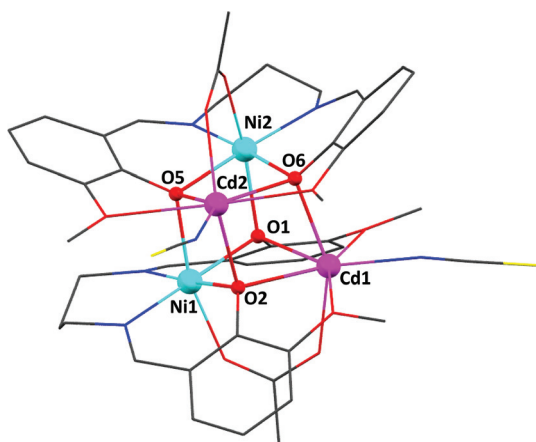


Fig. 1 Perspective view of complex 1 (unit A). Only the $\text{Cd}_2\text{Ni}_2\text{O}_4$ cubane core has been labelled. Hydrogen atoms have been omitted for clarity.

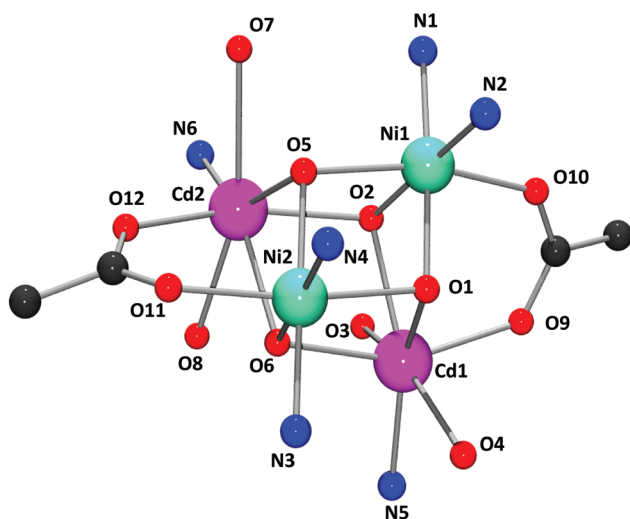


Fig. 2 Perspective view of the $\text{Cd}_2\text{Ni}_2\text{O}_4$ cubane core of unit A, along with the coordinating atoms and bridging acetate ligands.

Table 2 Selected bond lengths (Å) for complexes 1 (unit A) and 2

Complex	1	2	Complex	1	2
Cd(1)–O(1)	2.364(7)	2.238(2)	Ni(1)–O(2)	2.059(7)	—
Cd(1)–O(2)	2.390(6)	2.220(3)	Ni(1)–O(5)	2.190(7)	—
Cd(1)–O(3)	2.551(8)	2.581(4)	Ni(1)–O(10)	2.078(8)	—
Cd(1)–O(4)	2.507(9)	2.639(4)	Ni(2)–N(3)	2.040(9)	—
Cd(2)–O(6)	2.390(6)	2.232(3)	Ni(2)–O(5)	2.072(6)	—
Cd(2)–O(7)	2.584(8)	2.245(3)	Ni(2)–O(1)	2.177(7)	—
Cd(2)–O(8)	2.538(9)	2.543(3)	Ni(2)–N(4)	2.062(8)	—
Ni(1)–N(2)	2.059(9)	2.034(4)	Cd(1)–N(6)	—	2.194(6)
Ni(1)–N(1)	2.043(9)	2.036(4)	Cd(1)–N(7)	—	2.150(6)
Ni(1)–O(1)	2.085(6)	2.022(3)	Cd(2)–O(9)	—	2.669(3)
Ni(2)–O(11)	2.093(8)	2.107(4)	Cd(2)–S(7)	—	2.521(15)
Ni(2)–O(6)	2.062(7)	2.024(3)	Cd(2)–N(10)	—	2.175(4)
Cd(1)–O(6)	2.310(6)	—	Ni(1)–N(5)	—	2.141(4)
Cd(1)–O(9)	2.202(8)	—	Ni(1)–O(2)	—	2.011(4)
Cd(1)–N(5)	2.188(11)	—	Ni(1)–O(5)	—	2.112(3)
Cd(2)–O(2)	2.311(6)	—	Ni(2)–O(7)	—	2.021(3)
Cd(2)–O(5)	2.352(7)	—	Ni(2)–O(10)	—	2.100(3)
Cd(2)–O(12)	2.214(8)	—	Ni(2)–N(8)	—	2.031(4)
Cd(2)–N(6)	2.189(11)	—	Ni(2)–N(9)	—	2.025(4)

while each cadmium(II) resides in the outer O_4 compartment. Each nickel(II) centre is attached with two imine nitrogen atoms, [N(1) and N(2) for Ni(1); N(3) and N(4) for Ni(2)] and two phenoxo oxygen atoms, [O(1) and O(2) for Ni(1); O(5) and O(6) for Ni(2)] of the deprotonated Schiff base (L^1)²⁻ constituting the equatorial plane. The fifth coordination site of each nickel(II) centre is occupied by another phenoxo oxygen atom, [O(5) for Ni(1) and O(1) for Ni(2)], from the second deprotonated Schiff base. The octahedral geometries around both nickel(II) centres are fulfilled by the coordination of two oxygen atoms [O(10) for Ni(1), which bridge between Ni(1) and Cd(1), and O(11) for Ni(2), which bridges between Ni(2) and Cd(2)] from the two bridging acetates. For the Ni(1) centre, the deviation of the coordinating atoms, O(1), O(2), N(1) and N(2) in the basal plane from the mean plane passing through them are 0.002(6), 0.008(6), 0.008(9) and 0.002(8) Å, respectively. The deviation of Ni(1) from the same plane is $-0.020(13)$ Å. For the Ni(2) centre, the deviation of the coordinating atoms O(5), O(6), N(3) and N(4) in the basal plane from the mean plane passing through them are 0.007(6), $-0.014(6)$, $-0.013(9)$ and 0.006(8) Å, respectively. The deviation of Ni(2) from the same plane is 0.015(1) Å. Each cadmium(II) resides in outer O_4 compartment coordinated by two phenoxo [O(1) and O(2) for Cd(1); O(5) and O(6) for Cd(2)] and two methoxo oxygen atoms [O(3) and O(4) for Cd(1); O(7) and O(8) for Cd(2)]. The fifth coordination site in each cadmium(II) is occupied by two terminal N-bonded thiocyanates [N(5) for Cd(1) and N(6) for Cd(2)]. Phenoxo oxygen atom [O(6) for Cd(1) and O(2) for Cd(2)] coordinates in the sixth coordination site of each cadmium(II) centre axially. The pentagonal bipyramidal geometry around each cadmium(II) is completed by the coordination of oxygen atom [O(9) for Cd(1) and O(12) for Cd(2)] of the bridging acetate groups.

Cadmium(II) and nickel(II) centres in the cubane unit reside in an identical environment. Within the cubane unit, the Cd(1)–Ni(1), Ni(2)–Cd(2) and Ni(2)–Ni(1) distances are significantly



shorter (~ 3.32 Å) compared to the Cd(1)–Ni(2) and Cd(2)–Ni(1) distances (~ 3.44 Å), while the Cd(1)–Cd(2) distance is relatively longer (~ 3.69 Å). The cubane core has six faces. The Cd₂O₂ face is opposite the Ni₂O₂ face. The bridging angles M–O–M (M = metal) are comparable ($\sim 103^\circ$). The saturated six-membered chelate rings [Ni(1)–N(1)–C(9)–C(10)–C(11)–N(2)] and [Ni(2)–N(3)–C(31)–C(32)–C(33)–N(4)] have envelope conformations with the puckering parameters: $q = 0.523(14)$ Å; $\theta = 143.3(12)^\circ$; $\phi = 354(2)^\circ$ and $q = 0.503(13)$ Å; $\theta = 144.9(12)^\circ$; $\phi = 2(2)^\circ$, respectively.²⁵ The structure of unit **B** is similar to that of unit **A** and is described in the ESI.†

Complex **1** shows significant C–H \cdots π interactions. The hydrogen atom, H(9B), attached to carbon atom, C(9), is involved in an intra-molecular C–H \cdots π interaction with the phenyl ring [C(24)–C(25)–C(26)–C(27)–C(28)–C(29)]. Similarly the hydrogen atom, H(55B), attached to carbon atom, C(55), is involved in another intra-molecular C–H \cdots π interaction with the phenyl ring, C(68)–C(69)–C(70)–C(71)–C(72)–C(73). Another intra-molecular C–H \cdots π interaction is observed between the hydrogen atom, H(75A), attached to carbon atom, C(75A), with the phenyl ring, C(57)–C(58)–C(59)–C(60)–C(61)–C(62). The hydrogen atom, H(36), attached to carbon atom, C(36), shows another C–H \cdots π interaction with the phenyl ring, C(68)–C(69)–C(70)–C(71)–C(72)–C(73). These C–H \cdots π interactions are shown in Fig. S2 {ESI†}. The hydrogen atom, H(80), attached to carbon atom, C(80), shows an inter-molecular C–H \cdots π interaction with a symmetry related ($x, 1 + y, z$) phenyl ring [C(2)–C(3)–C(4)–C(5)–C(6)–C(7)], as shown in Fig. S3.† Another C–H \cdots π inter-molecular interaction is observed between the hydrogen atom, H(49), attached to carbon atom, C(49), with the symmetry related ($-1 + x, y, z$) phenyl ring [C(24)–C(25)–C(26)–C(27)–C(28)–C(29)], as shown in Fig. S4.† The hydrogen atom, H(14), attached to carbon atom, C(14), is involved in another inter-molecular C–H \cdots π interaction with the symmetry related ($1 + x, -1 + y, z$) phenyl ring [C(57)–C(58)–C(59)–C(60)–C(61)–C(62)], as shown in Fig. S5.† The details of the geometric features of the C–H \cdots π interactions are given in Table 3.

$[(\text{DMSO})_2\text{NiL}^2\text{Cd}(\text{NCS})(\mu_{1,3}\text{-SCN})\text{Ni}(\text{DMSO})\text{L}^2\text{Cd}(\text{NCS})_2]$ (**2**). The X-ray crystal structure determination revealed that complex **2** crystallizes in the monoclinic space group $P2_1$. The asymmetric unit contains a tetranuclear neutral species, $[(\text{DMSO})_2\text{NiL}^2\text{Cd}(\text{NCS})(\mu_{1,3}\text{-SCN})\text{Ni}(\text{DMSO})\text{L}^2\text{Cd}(\text{NCS})_2]$ (Fig. 3). The metal centres and coordinating atoms and bridging thiocyanate are shown in Fig. 4. The tetranuclear moiety consists of two pseudo-dinuclear units: **C** $[\text{Ni}(\text{DMSO})\text{L}^2\text{Cd}(\text{NCS})_2]$ (Fig. 5a) and **D** $[(\text{DMSO})_2\text{NiL}^2\text{Cd}(\text{NCS})]^+$ (Fig. 5b), which are joined through a single end-to-end thiocyanate bridge. The important bond lengths and bond angles are gathered in Tables 2 and S2 {ESI†}, respectively. In the dimeric unit **C**, the Ni(1) and Cd(1) centres occupy the inner N₂O₂ and outer O₄ sites, respectively. The Ni(1) and Cd(1) centres are bridged by two phenoxo oxygen atoms, O(1) and O(2), with a Ni(1) \cdots Cd(1) distance of $3.324(8)$ Å. Both Ni(1) and Cd(1) have pseudo-octahedral geometries. Ni(1) is coordinated by two imine nitrogen atoms, N(1) and N(2), and two phenoxo oxygen atoms, O(1) and O(2), of one deprotonated potential octadentate Schiff base ligand (L²)²⁻, which constitute the equatorial plane. The

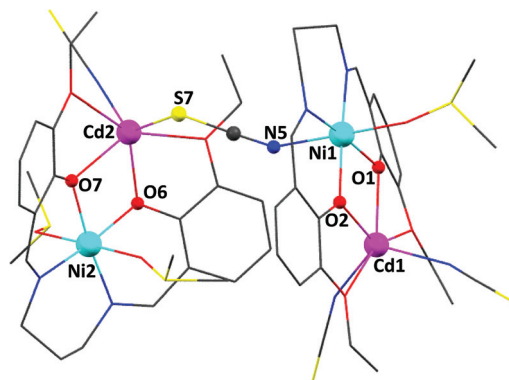


Fig. 3 Perspective view of complex **2** with selective atom numbering. Hydrogen atoms have been omitted for clarity.

Table 3 Geometric features (distances in Å and angles in $^\circ$) of the C–H \cdots π interactions obtained for complexes **1** and **2**

Complex	X–H \cdots Cg(Ring)	H \cdots Cg (Å)	C–H \cdots Cg ($^\circ$)	C \cdots Cg (Å)
1	C(14)–H(14) \cdots Cg(20) ^a	2.83	135	3.549(15)
	C(36)–H(36) \cdots Cg(21)	2.84	129	3.495(15)
	C(49)–H(49) \cdots Cg(10) ^b	2.90	126	3.527(15)
	C(9)–H(9B) \cdots Cg(10)	2.93	151	3.807(14)
	C(55)–H(55B) \cdots Cg(21)	2.97	148	3.830(15)
	C(75)–H(75A) \cdots Cg(20)	2.97	150	3.842(15)
	C(80)–H(80) \cdots Cg(8) ^c	2.86	134	3.566(15)
	2	C(48)–H(48A) \cdots Cg(14) ^d	2.91	133
C(41)–H(41) \cdots Cg(13) ^e		2.85	149	3.672(7)
C(29)–H(29) \cdots Cg(13)		2.98	153	3.829(6)

Symmetry transformations: ^a = $1 + x, -1 + y, z$; ^b = $-1 + x, y, z$; ^c = $x, 1 + y, z$; ^d = $-x, 2 - y, -z$; ^e = $\frac{1}{2} + x, 3/2 - y, \frac{1}{2} + z$. Cg(8) = centre of gravity of the ring [C(2)–C(3)–C(4)–C(5)–C(6)–C(7)]; Cg(10) = centre of gravity of the ring [C(24)–C(25)–C(26)–C(27)–C(28)–C(29)]; Cg(20) = centre of gravity of the ring [C(57)–C(58)–C(59)–C(60)–C(61)–C(62)]; Cg(21) = centre of gravity of the ring [C(68)–C(69)–C(70)–C(71)–C(72)–C(73)] for complex **1** and Cg(13) = centre of gravity of the ring [C(7)–C(8)–C(9)–C(10)–C(11)–C(12)]; Cg(14) = centre of gravity of the ring [C(18)–C(19)–C(20)–C(21)–C(22)–C(23)]; for complex **2**.



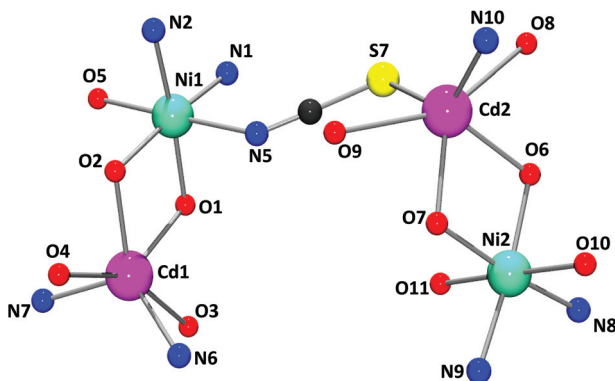


Fig. 4 Perspective view of the tetranuclear unit of complex 2 with selective atom numbering. Only the metal centres with coordinating atoms and the bridging thiocyanate are shown for clarity.

fifth coordination site is occupied by one oxygen atom, O(5) from a DMSO molecule, whereas, the sixth coordination site of Ni(1) is occupied by a nitrogen atom, N(5), of an end-to-end bridged thiocyanate, which links the two pseudo-dinuclear units C and D. On the other hand, Cd(1) is coordinated by two phenoxo oxygen atoms, O(1) and O(2), and two ethoxy oxygen atoms, O(3) and O(4), which constitute the equatorial plane. The fifth and sixth coordination sites are occupied by two nitrogen atoms, N(6) and N(7), from two terminal thiocyanates. The bridging angles Ni(1)–O(1)–Cd(1) and Ni(1)–O(2)–Cd(1) are 102.4(1)° and 103.4(1)°, respectively. The angles O(1)–Ni(1)–O(2) and O(1)–Cd(1)–O(2) are 81.6(1)° and 72.5(1)°, respectively. The saturated six-membered chelate ring [Ni(1)–N(1)–C(14)–C(15)–C(16)–N(2)] has envelope conformations with the puckering parameters: $q = 0.490(9)$ Å; $\theta = 132(8)^\circ$; $\phi = 347.7(11)^\circ$.²⁵

The dinuclear unit D has a similar structure to unit C. Unit D contains nickel(II) and cadmium(II) centres, Ni(2), Cd(2), which are bridged by two phenoxo oxygen atoms, O(6) and O(7), with a Ni(2)⋯Cd(2) distance of 3.300(6) Å. Ni(2) is

coordinated by two imine nitrogen atoms, N(8) and N(9), and two phenoxo oxygen atoms, O(6) and O(7), of the deprotonated potential octadentate Schiff base ligand (L²⁻). The remaining two coordination sites of Ni(2) are occupied by two oxygen atoms, O(10) and O(11), from two DMSO molecules. On the other hand, Cd(2) is coordinated by two phenoxo oxygen atoms, O(6) and O(7), two ethoxy oxygen atoms, O(8) and O(9), and one nitrogen atom, N(10), from one terminal thiocyanate. The sixth coordination site of Cd(2) is occupied by a sulphur atom, S(7), of an end-to-end bridged thiocyanate, which links the two pseudo-dinuclear units C and D. The saturated six-membered chelate ring [Ni(2)–N(8)–C(35)–C(36)–C(37)–N(9)] has an envelope conformation with the puckering parameters: $q = 0.528(6)$ Å; $\theta = 144.1(5)^\circ$; $\phi = 3.5(10)^\circ$.²⁵ The Ni(1)⋯Cd(2) separation in the tetranuclear unit is 6.42(7) Å.

Complex 2 shows three C–H⋯π interactions. The hydrogen atom, H(48A), attached to carbon atom, C(48), is involved in an inter-molecular (between two C units) C–H⋯π interaction with a symmetry related ($-x, 2 - y, -z$) phenyl ring [C(18)–C(19)–C(20)–C(21)–C(22)–C(23)] to form a supramolecular dimer. One intra-molecular C–H⋯π interaction is observed between two different subunits of the complex having different symmetries. The hydrogen atom, H(29), attached to carbon atom, C(29), is involved in an intra-molecular (between the C and D units) C–H⋯π interaction with a phenyl ring [C(7)–C(8)–C(9)–C(10)–C(11)–C(12)]. These interactions are shown in Fig. S6 {ESI†}. The hydrogen atom, H(41), attached to carbon atom, C(41), is involved in another inter-molecular (between the C and D units) C–H⋯π interaction with the symmetry related ($1/2 + x, 3/2 - y, 1/2 + z$) phenyl ring, C(7)–C(8)–C(9)–C(10)–C(11)–C(12) to form another supramolecular dimer, as shown in Fig. S7.† The details of the geometric features of the C–H⋯π interactions are given in Table 3. In addition to these C–H⋯π interactions, complex 2 also shows S⋯π interactions. A remarkable S⋯π interaction is observed between the lone pair of disordered sulphur atoms with the C–N and C–S orbitals of the thiocyanate [N(7)–C(2)–S(9)], attached with Cd(1). In this

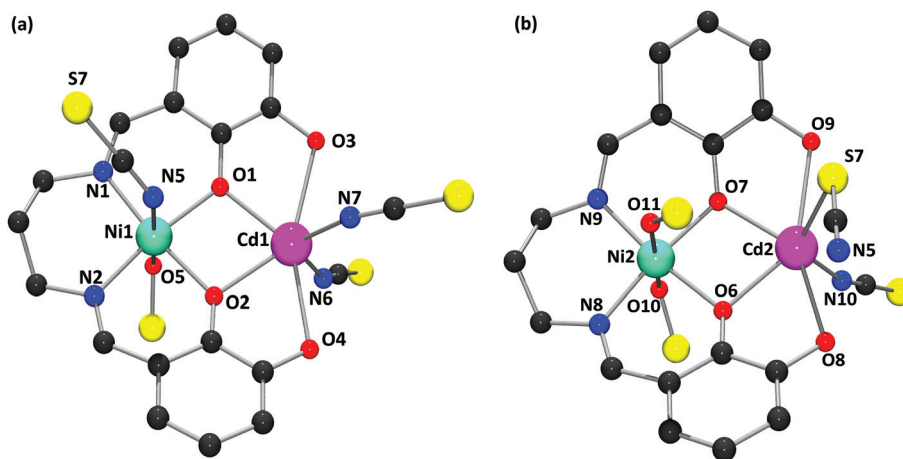


Fig. 5 Perspective views of the pseudo-dinuclear units [(a) unit C; (b) unit D] of complex 2 with selective atom numbering. Hydrogen atoms and the methyl groups of DMSO and ethyl groups have been omitted for clarity.



S... π interaction, DMSO acts as an electron-rich moiety (electron donor) while thiocyanate acts as an electron acceptor in spite of its anionic character. Details of this S... π interaction are explored in the theoretical part of this study below.

Theoretical study

The theoretical study was divided into two parts. Firstly, the surprisingly different architecture observed in the solid state of the two complexes was rationalized taking into account that the only difference between the H₂salen-type Schiff bases is the alkoxy substituent (methoxy or ethoxy). The supramolecular assemblies observed in the crystal packing of complexes **1** and **2** has also been analyzed.

As described above, in the cubane assembly, each cadmium(II) centre is heptacoordinated, being coordinated by two phenoxo and two methoxy oxygen atoms, with the fifth coordination site in the equatorial plane being occupied by a terminal N-bonded thiocyanate. This coordination mode, which is required for the formation of the cubane, is only possible for the methoxy substituted ligand, since the steric hindrance of both ethoxy groups prevents the formation of the heptacoordinated cadmium atom (only four atoms in the equatorial plane) in the L² ligand (Fig. 6). As a matter of fact, the cadmium ions are hexacoordinated in compound **2**, giving rise to a totally different complex.

In the crystal structure of **2**, the nickel(II) and cadmium(II) metal centres are bridged by only one NCS ligand, in which the nickel(II) of the C unit is connected with the cadmium(II) of the D unit through the NCS bridge. On the other hand, the nickel(II) of the D unit is not coordinated by the NCS ligand from the C unit but is rather coordinated by two DMSO molecules. One of both DMSO molecules establishes a non-covalent S... π interaction, which is studied below. The coordination of this DMSO molecule prevents the formation of the doubly NCS-bridged structure. We analysed this issue by means of DFT calculations. We optimized the hypothetical double bridge structure and compared the formation energy with the one observed experimentally. The results are shown in Fig. 7, and it can be observed that the dissymmetric assembly

observed experimentally is 14.4 kcal mol⁻¹ more stable than the doubly bridged structure. A likely explanation is that both assemblies present the same number of metal–ligand coordination bonds but the experimentally observed complex has some additional stabilization from the intra-molecular non-covalent interactions established by the DMSO molecule.

The second part of the theoretical study is devoted to analysing the unconventional non-covalent interactions found in the crystal packing of complexes **1** and **2**, focusing our attention to the C–H...H–C interactions and the remarkable S... π interactions. In **1**, we analyzed the interesting C–H...H–C interactions observed between the organic ligand and acetate co-ligand in the solid-state structure. In **2**, we analyzed the remarkable and unprecedented S... π interaction observed in the solid-state structure by using several computational tools, such as NBO and AIM analyses. The importance of the C–H...H–C interaction was highlighted by Alvarez and co-workers.²⁶ Dihydrogen contacts in alkanes are among the weakest inter-molecular interactions (\sim –0.4 kcal mol⁻¹ for the methane dimer); however, they are cumulative, resulting in larger dimerization energies in some cases (e.g. long open chains). For example, the complexation energy computed for the dimer of *n*-hexane is –4.5 kcal mol⁻¹.²⁶

It is also worthy to emphasize that supramolecular chemists usually analyze the conventional non-covalent interactions in the X-ray structures (shorter than the sum of the van der Waals radii), ignoring longer interactions in the crystal structures. Evidently, this is a simplistic view of the crystal packing since interactions slightly longer also have an impact. Bearing this in mind, we studied the interesting crystal packing observed in **1** (Fig. 8a), where 2D planes are found in the solid state governed basically by the long range dispersion and C–H...H–C interactions. Curiously, the tetranuclear complexes are arranged in such a way that four methyl groups belonging to the acetate ligands of four different complexes point to the same void space (Fig. 8a). In a similar way, four methyl groups of the methoxy substituent of the aromatic ligand also point to the void space of the 2D plane (the blue lines in Fig. 8a). Moreover, the formation of the 2D plane is due to the van der

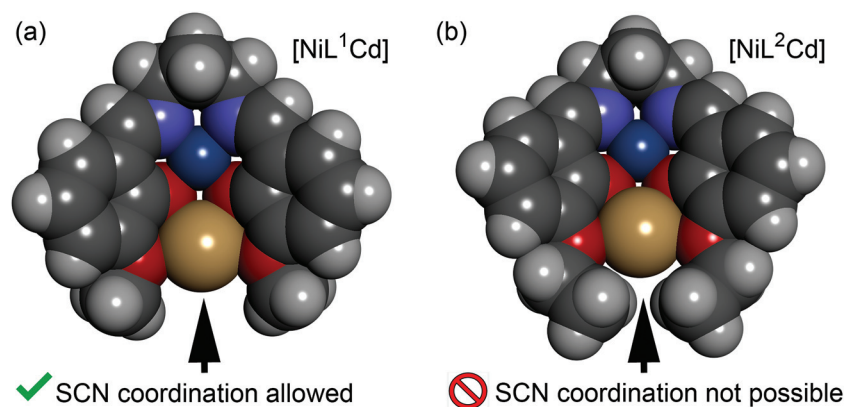


Fig. 6 CPK representation of the nickel(II)/cadmium(II) complexes with L¹ (a) and L² (b).



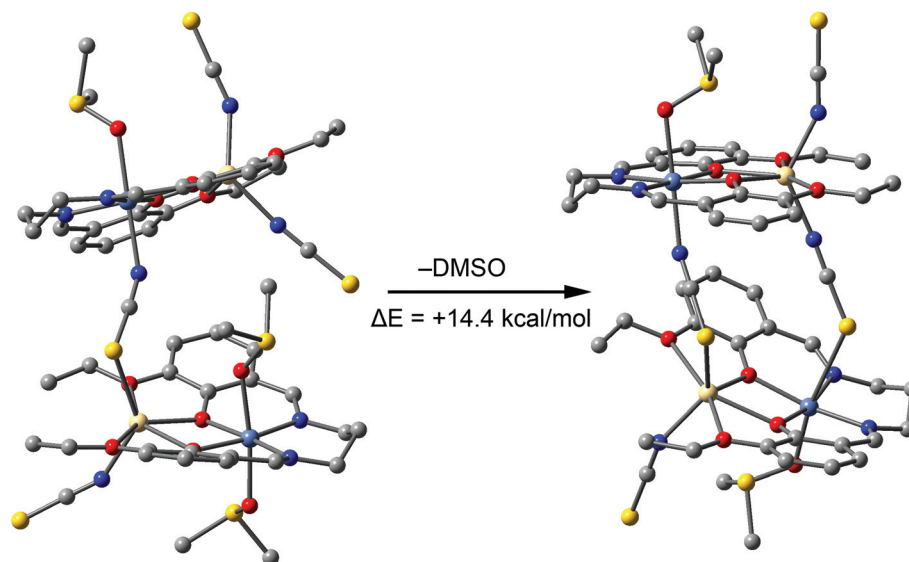


Fig. 7 Optimized geometries of the dissymmetric assembly of complex 2 (left) and the hypothetical doubly bridged assembly. The formation energy is also indicated.

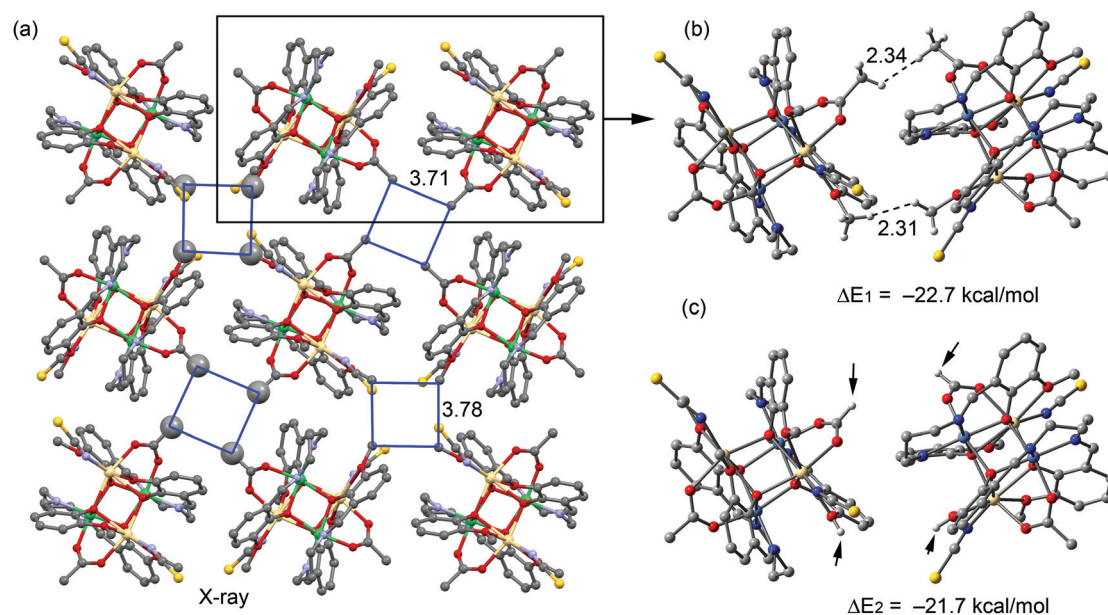


Fig. 8 (a) Partial view of the X-ray solid-state structure of 1. H atoms have been omitted for clarity. Some $-\text{CH}_3$ groups are highlighted using a higher ball radius in the ball and stick representation; (b, c) theoretical models used to evaluate the $\text{C}-\text{H}\cdots\text{H}-\text{C}$ interaction. Only the H atoms of the interacting methyl groups are shown. Distances in Å.

Waals dispersion interactions between the bulk of the molecules. In order to evaluate the contribution of the $\text{C}-\text{H}\cdots\text{H}-\text{C}$ interactions involving the methyl groups, we used a theoretical model consisting of one dimer taken from this 2D plane, as shown in Fig. 8b (using the crystallographic coordinates). We computed its interaction energy, which is $\Delta E_1 = -22.7 \text{ kcal mol}^{-1}$ and which could be attributed to the contribution of two $\text{CH}_3\cdots\text{H}_3\text{C}$ interactions (in one, the $-\text{CH}_3$ groups are from the acetate and in the other they are from the

methoxy group, see dashed lines) along with the long range interactions. In an effort to estimate the contribution of the $\text{C}-\text{H}\cdots\text{H}-\text{C}$ interaction, we computed an additional model, where the methyl groups were changed by hydrogen atoms (the small arrows in Fig. 8c). As a result, the interaction is modestly reduced to $\Delta E_2 = -21.7 \text{ kcal mol}^{-1}$, which is the contribution of the van der Waals dispersion interactions between the bulk of the two molecules. The difference between both interaction energies ($\Delta E_1 - \Delta E_2 = -1 \text{ kcal mol}^{-1}$) is the contri-



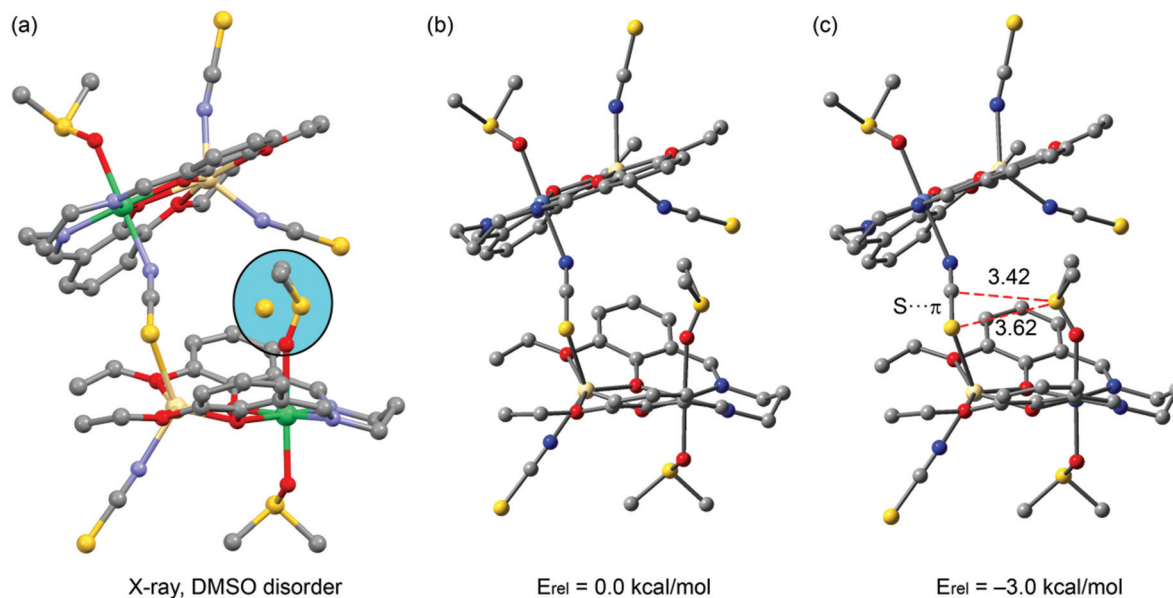


Fig. 9 (a) X-ray solid-state structure of **2**; (b, c) theoretical models used to evaluate the S... π interaction. H atoms omitted for clarity. Distances in Å.

tribution of the CH₃...H₃C interactions. This value is in good agreement with the reported calculations of the methane dimer.

In complex **2**, we analyzed the disordered DMSO molecule observed in its solid-state X-ray structure (Fig. 9a). The sulphur atom is disordered in two positions and in one of both the sulphur atom is in contact with the central carbon atom of the bridging thiocyanate. This interaction can be viewed as a contact between the electron-rich sulphur atom with the π -system of the thiocyanate. In fact, we previously demonstrated²⁷ the ability of pseudo-halide ligands (thiocyanate and selenocyanate) to interact with either H bonds (via the ending atoms) or lone pair donors (via the central carbon atom). We evaluated energetically the difference between the two disordered positions at the M06-2X/def2-TZVP level of theory, and the one with the S... π interaction was more stable by 3 kcal mol⁻¹, which can be attributed to this interaction.

At this point, in order to investigate the S... π interaction from an orbital point of view, we performed Natural Bond Orbital (NBO) calculations in the model of Fig. 9c, focusing our attention on the second-order perturbation analysis, which is very useful to study donor-acceptor interactions.²⁸ Interestingly, we found that the lone pair (lp orbital) of the sulphur atom of DMSO interacts with the C-N and C-S antibonding orbitals (BD*) of the thiocyanate, with a concomitant second-order stabilization energy of $E^{(2)} = 0.98$ kcal mol⁻¹ for the LP(S) \rightarrow BD*(CN) interaction and $E^{(2)} = 0.05$ kcal mol⁻¹ for the LP(S) \rightarrow BD*(CS) orbital interaction. Therefore, the global orbital stabilization energy that can be attributed to the S... π interaction is approximately 1.03 kcal mol⁻¹ in the dimer (~33% of the total interaction energy). This analysis confirms both the existence of the interaction and also that the DMSO is acting as an electron-rich moiety (electron donor) and the thiocyanate moiety as an electron acceptor, in spite of its anionic

character. This is likely due to the double coordination of the thiocyanate to two divalent metal centres [nickel(II) and cadmium(II)], which drastically increases the π -acidity of the ligand. We also computed the AIM analysis of the model of complex **2** shown in Fig. 9c in order to investigate the distribution of the critical points, which is shown in Fig. 10. A bond critical point and bond path that connects the sulphur atom of

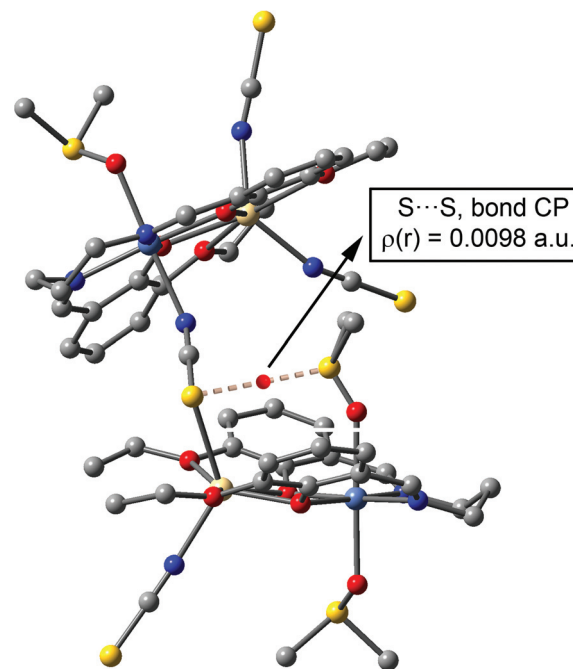


Fig. 10 Representation of the bond critical point (red sphere) and bond path connecting the S atom of DMSO to the S atom of the pseudo-halide ligand (dashed line). H atoms omitted for clarity.



Table 4 Photophysical data for complexes **1** and **2**

Complex	Absorption (nm)	Emission (nm)	A ₁ (%)	τ ₁ (ns)	A ₂ (%)	τ ₂ (ns)	Γ _{avg} (ns)	χ ²
1	375	453	33.64	6.94	66.36	30.12	27.7	1.1098
2	378	456	33.64	7.21	66.36	29.20	26.8	1.1481

the DMSO with the sulphur atom of the pseudo-halide ligand can be observed, thus confirming the S...π interaction. The rest of the critical points and bond paths in this assembly have been omitted for clarity.

IR and electronic spectra and the photophysical study

In the IR spectra of both complexes, distinct bands due to the azomethine (C=N) groups within the range of 1626–1632 cm⁻¹ are routinely noticed.²⁹ A strong band at 2067 cm⁻¹ indicates the presence of terminal N-bonded thiocyanate in complex **1**. There are two strong bands at 2034 and 2068 cm⁻¹ in complex **2**, indicating the presence of terminal (N-bonded) and end-to-end bridged thiocyanate³⁰ respectively, which are also evident from the crystal structure determination.

As cadmium(II) is a d¹⁰ system, no d–d bands are observed due to cadmium(II). The intense absorption bands at shorter wavelengths (~375 nm) may be assigned to metal-to-ligand charge-transfer bands (MLCT) due to both the nickel(II) and cadmium(II) centres.^{31a} In the visible range, two absorption bands around 670 nm and 900 nm are observed. These bands may be assigned as ³T_{1g}(F) ← ³A_{2g}(F) and ³T_{2g}(F) ← ³A_{2g}(F), respectively.^{31b,c} The higher energy d–d band, ³T_{1g}(P) ← ³A_{2g}(F), expected for octahedral nickel(II), cannot be observed as it is obscured by strong charge-transfer transitions (~375 nm).^{31c} The bands at ~270 nm may be assigned as ligand centred π* ← π transitions.³²

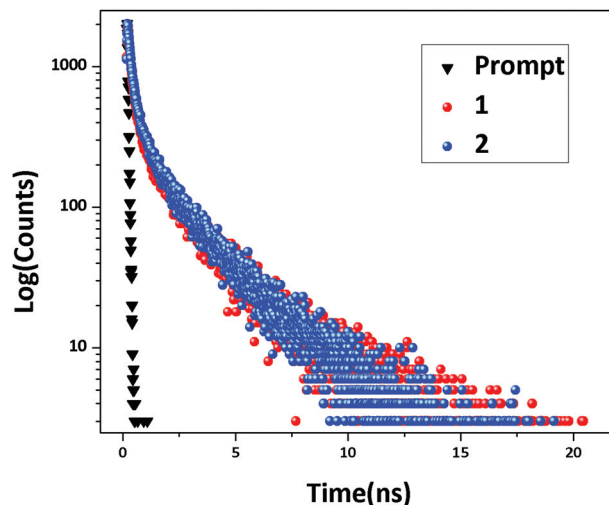
Both complexes exhibit fluorescence in DMSO medium. The fluorescence data are listed in Table 4 (without solvent correction). These are assigned as intra-ligand (π* ← π) fluorescence.³² The mean lifetimes (Γ_{avg}) of the excited states are 27.7 ns (for **1**) and 26.8 ns (for **2**) at room temperature (Table 4). Decay profiles (Fig. 11) were fitted to a multi-exponential model:

$$I(t) = \sum_i \alpha_i e^{-\frac{t}{\tau_i}}$$

where bi-exponential functions were used to fit the emission of all the complexes and to obtain χ² close to 1. The intensity-averaged lifetimes (Γ_{avg}) were determined using the following equation:

$$\langle \tau \rangle = \frac{\sum_i \alpha_i \tau_i^2}{\sum_i \alpha_i \tau_i}$$

where α_i and τ_i are the pre-exponential factor and excited-state luminescence decay time associated with the *i*-th component, respectively.

**Fig. 11** Lifetime decay profile of complexes **1** and **2**.

PXRD

The experimental PXRD patterns of the bulk products were in good agreement with the simulated XRD patterns from the single crystal X-ray diffraction results, indicating consistency of the bulk sample. The simulated patterns of the complexes were calculated from the single crystal structural data (Cif) using the CCDC Mercury software.

Concluding remarks

Synthesis and X-ray characterization of two new hetero-tetra-nuclear nickel(II)–cadmium(II) complexes with Schiff base ligands have been described in this paper. Between the complexes, complex **1** was a phenoxo-bridged cubane and complex **2** was a linear tetramer with Ni₂Cd₂O₄ cores. The linear tetramer consisted of two pseudo-dinuclear units, joined through a single end-to-end thiocyanate bridge. The solid-state structure of complex **1** showed the participation of the organic ligand and acetate group in C–H...H–C interactions, which were evaluated energetically by means of DFT calculations. Moreover, the disorder of the DMSO molecule was studied in complex **2**, since one of the disordered DMSO molecules presented an unprecedented S...π interaction, where the π-system was provided by the thiocyanate. This unexpected interaction was evidenced by means of several computational tools, such as NBO and AIM analyses.



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Notes and references

- (a) A. Hazari, L. K. Das, A. Bauzá, A. Frontera and A. Ghosh, *Dalton Trans.*, 2016, **45**, 5730–5740; (b) P. Seth, S. Giri and A. Ghosh, *Dalton Trans.*, 2015, **44**, 12863–12870; (c) B. J. Kennedy and K. S. Murray, *Inorg. Chem.*, 1985, **24**, 1552–1557; (d) A. van den Bergen, K. S. Murray, B. O. West and A. N. Buckley, *J. Chem. Soc. A*, 1969, 2051–2060; (e) E. Uhlig, *Coord. Chem. Rev.*, 1973, **10**, 227–264; (f) S. Yamada, *Coord. Chem. Rev.*, 1999, **190**, 537–555; (g) P. Mukherjee, C. Biswas, M. G. B. Drew and A. Ghosh, *Polyhedron*, 2007, **26**, 3121–3128; (h) L. S. Felices, E. C. Escudero-Adán, J. Benet-Buchholz and A. W. Kleij, *Inorg. Chem.*, 2009, **48**, 846–853; (i) S. A. Fairhurst, D. L. Hughes, G. J. Leigh, J. R. Sanders and J. Weisner, *J. Chem. Soc., Dalton Trans.*, 1994, 2591–2598; (j) R. M. Clarke and T. Storr, *Dalton Trans.*, 2014, **43**, 9380–9391; (k) X. Yanga, R. A. Jonesb and S. Huanga, *Coord. Chem. Rev.*, 2014, **273–274**, 63–75.
- (a) N. Sarkar, P. K. Bhaumik and S. Chattopadhyay, *Polyhedron*, 2016, **115**, 37–46; (b) P. Bhowmik, H. P. Nayek, M. Corbella, N. Aliaga-Alcalde and S. Chattopadhyay, *Dalton Trans.*, 2011, **40**, 7916–7926; (c) E. López-Torres and M. A. Mendiola, *Dalton Trans.*, 2009, 7639–7647; (d) M. Sutradhar, T. Roy Barman, M. G. B. Drew and E. Rentschler, *J. Mol. Struct.*, 2013, **1041**, 44–49; (e) D. A. Rotsch, K. M. Reinig, E. M. Weis, A. B. Taylor, C. L. Barnesa and S. S. Jurisson, *Dalton Trans.*, 2013, **42**, 11614–11625; (f) S. Chandra and R. Kumar, *Transition Met. Chem.*, 2004, **29**, 269–275; (g) S. A. Schuetz, C. M. Silvernail, C. D. Incarvito, A. L. Rheingold, J. L. Clark, V. W. Day and J. A. Belot, *Inorg. Chem.*, 2004, **43**, 6203–6214; (h) S. Mohebbi and M. Abdi, *J. Coord. Chem.*, 2008, **61**, 3410–3419; (i) A. Ray, S. Banerjee, G. M. Rosair, V. Gramlich and S. Mitra, *Struct. Chem.*, 2008, **19**, 459–465; (j) D. Mandal, P. B. Chatterjee, S. Bhattacharya, K.-Y. Choi, R. Clérac and M. Chaudhury, *Inorg. Chem.*, 2009, **48**, 1826–1835; (k) A. Vlad, M.-F. Zaltariov, S. Shova, M. Cazacu, M. Avadanei, A. Soroceanu and P. Samoila, *Polyhedron*, 2016, **115**, 76–85.
- (a) S. Bhattacharya and S. Mohanta, *Inorg. Chim. Acta*, 2015, **432**, 169–175; (b) P. A. Vigato and S. Tamburini, *Coord. Chem. Rev.*, 2008, **252**, 1871–1995; (c) L. K. Das, C. J. Gómez-García, M. G. B. Drew and A. Ghosh, *Polyhedron*, 2015, **87**, 311–320; (d) P. Chakraborty and S. Mohanta, *Polyhedron*, 2015, **87**, 98–108.
- (a) S. Bhattacharya, A. Jana and S. Mohanta, *CrystEngComm*, 2013, **15**, 10374–10382; (b) A. Biswas, L. Mandal, S. Mondal, C. R. Lucas and S. Mohanta, *CrystEngComm*, 2013, **15**, 5888–5897; (c) P. Bhowmik, H. P. Nayek, M. Corbella, N. Aliaga-Alcalde and S. Chattopadhyay, *Dalton Trans.*, 2011, **40**, 7916–7926; (d) P. Bhowmik, S. Jana, P. P. Jana, K. Harms and S. Chattopadhyay, *Inorg. Chim. Acta*, 2012, **390**, 53–60; (e) P. Bhowmik, S. Jana, P. P. Jana, K. Harms and S. Chattopadhyay, *Inorg. Chem. Commun.*, 2012, **18**, 50–56; (f) P. Bhowmik, S. Jana and S. Chattopadhyay, *Polyhedron*, 2012, **44**, 11–17; (g) P. Bhowmik, K. Harms and S. Chattopadhyay, *Polyhedron*, 2013, **49**, 113–120; (h) P. Bhowmik, N. Aliaga-Alcalde, V. Gómez, M. Corbella and S. Chattopadhyay, *Polyhedron*, 2013, **49**, 269–276; (i) P. Bhowmik, S. Chatterjee and S. Chattopadhyay, *Polyhedron*, 2013, **63**, 214–221.
- (a) K. Bhattacharya, M. Maity, S. Md T. Abtab, M. C. Majee and M. Chaudhury, *Inorg. Chem.*, 2013, **52**, 9597–9605; (b) A. Jana, R. Koner, T. Weyhermueller, P. Lemoine, M. Ghosh and S. Mohanta, *Inorg. Chim. Acta*, 2011, **375**, 263–270.
- (a) E. Evangelio, N. P. Rath and L. M. Mirica, *Dalton Trans.*, 2012, **41**, 8010–8021; (b) X. Gong, Y.-Y. Ge, M. Fang, Z.-G. Gu, S.-R. Zheng, W.-S. Li, S.-J. Hu, S.-B. Lib and Y.-P. Cai, *CrystEngComm*, 2011, **13**, 6911–6915.
- (a) S. Sasmal, S. Hazra, P. Kundu, S. Majumder, N. Aliaga-Alcalde, E. Ruiz and S. Mohanta, *Inorg. Chem.*, 2010, **49**, 9517–9526; (b) S. Hazra, R. Koner, P. Lemoine, E. C. Sañudo and S. Mohanta, *Eur. J. Inorg. Chem.*, 2009, 3458–3466; (c) J. P. Wikstrom, A. Y. Nazarenko, W. M. Reiff and E. V. Rybak-Akimova, *Inorg. Chim. Acta*, 2007, **360**, 3733–3740; (d) B. Biswas, U. Pieper, T. Weyhermuller and P. Chaudhuri, *Inorg. Chem.*, 2009, **48**, 6781–6793; (e) J. M. Clemente-Juan, B. Chansou, B. Donnadiou and J. Tuchagues, *Inorg. Chem.*, 2000, **39**, 5515–5519; (f) S.-Y. Zhang, W.-Q. Chen, B. Hu, Y.-M. Chen, W. Li and Y. Li, *Inorg. Chem. Commun.*, 2012, **16**, 74–77; (g) S.-H. Zhang, N. Li, C.-M. Ge, C. Fenga and L.-F. Ma, *Dalton Trans.*, 2011, **40**, 3000–3007.
- (a) H. Y. Kwak, D. W. Ryu, H. C. Kim, E. K. Koh, B. K. Chod and C. S. Hong, *Dalton Trans.*, 2009, 1954–1961; (b) H. Chen, C.-B. Ma, D.-Q. Yuan, M.-Q. Hu, H.-M. Wen, Q.-T. Liu and C.-N. Chen, *Inorg. Chem.*, 2011, **50**, 10342–10352.
- (a) S. Biswas, C. J. Gómez-García, J. M. Clemente-Juan, S. Benmansour and A. Ghosh, *Inorg. Chem.*, 2014, **53**, 2441–2449; (b) L. K. Das and A. Ghosh, *CrystEngComm*, 2013, **15**, 9444–9456; (c) S. L. Roderick and B. W. Matthews, *Biochemistry*, 1993, **32**, 3907–3912; (d) K. Ghosh, S. Roy, A. Ghosh, A. Banerjee, A. Bauzá, A. Frontera and S. Chattopadhyay, *Polyhedron*, 2016, **112**, 6–17.
- (a) J. Chen, J. Tang, J. Zhou, L. Zhang, G. Chen and D. Tang, *Anal. Chim. Acta*, 2014, **810**, 10–16; (b) K. Mahata and M. Schmittel, *J. Am. Chem. Soc.*, 2009, **131**, 16544–16554.



- 11 (a) Y.-S. Yang, Y.-P. Yang, M. Liu, Q.-M. Qiu, Q.-H. Jin, J.-J. Sun, H. Chen, Y.-C. Dai and Q.-X. Meng, *Polyhedron*, 2015, **35**, 912–917; (b) Y. Shen, N. Ma, L. Wu and H.-H. Song, *Inorg. Chim. Acta*, 2015, **429**, 51–60; (c) A. Bhattacharyya, P. K. Bhaumik, P. P. Jana and S. Chattopadhyay, *Polyhedron*, 2014, **34**, 40–45; (d) S. Carboni, C. Gennari, L. Pignataro and U. Piarulli, *Dalton Trans.*, 2011, **40**, 4355–4373.
- 12 (a) C. A. Hunter and J. K. M. Sanders, *J. Am. Chem. Soc.*, 1990, **112**, 5525–5534; (b) S. K. Burley and G. A. Petsko, *Science*, 1985, **229**, 23–28; (c) K. S. Kim, P. Tarakeshwar and J. Y. Lee, *Chem. Rev.*, 2000, **100**, 4145–4185; (d) J. C. Ma and D. A. Dougherty, *Chem. Rev.*, 1997, **97**, 1303–1324; (e) K. S. Kim, J. Y. Lee, S. J. Lee, T.-K. Ha and D. H. Kim, *J. Am. Chem. Soc.*, 1994, **116**, 7399–7400; (f) M. Nishio, M. Hirota and Y. Umezawa, in *The C–H/p Interaction: Evidence, Nature, Consequences*, Wiley-VCH, New York, 1998; (g) M. Nishio, *CrystEngComm*, 2004, **6**, 130–156; (h) M. Egli and S. Sarkhel, *Acc. Chem. Res.*, 2007, **40**, 197–205; (i) T. J. Mooibroek, P. Gamez and J. Reedijk, *CrystEngComm*, 2008, **10**, 1501–1515; (j) J. Ran and P. Hobza, *J. Chem. Theory Comput.*, 2009, **5**, 1180–1185; (k) M. Barceló-Oliver, C. Estarellas, A. García-Raso, A. Terrón, A. Frontera, D. Quiñonero, E. Molins and P. M. Deyà, *CrystEngComm*, 2010, **12**, 362–365; (l) D. Quiñonero, C. Garau, C. Rotger, A. Frontera, P. Ballester, A. Costa and P. M. Deyà, *Angew. Chem., Int. Ed.*, 2002, **41**, 3389–3392.
- 13 (a) D. Paolantoni, J. Rubio-Magnieto, S. Cantel, J. Martinez, P. Dumy, M. Surin and S. Ulrich, *Chem. Commun.*, 2014, **50**, 14257; (b) S. Jana, K. Harms and S. Chattopadhyay, *J. Coord. Chem.*, 2014, **67**, 2954–2966; (c) C. Estarellas, A. Frontera, D. Quiñonero and P. M. Deyà, *Angew. Chem., Int. Ed.*, 2011, **50**, 415.
- 14 (a) S. Naskar, D. Mishra, R. J. Butcher and S. K. Chattopadhyay, *Polyhedron*, 2007, **26**, 3703–3714; (b) L. M. Salonen, M. Ellermann and F. Diederich, *Angew. Chem., Int. Ed.*, 2011, **50**, 4808; (c) J. Xiao, P. Broz, A. W. Puri, E. Deu, M. Morell, D. M. Monack and M. Bogyo, *J. Am. Chem. Soc.*, 2013, **135**, 9130–9138; (d) A. Bauzá, D. Quiñonero, P. M. Deyà and A. Frontera, *Chem. – Asian J.*, 2013, **8**, 2708–2713.
- 15 (a) A. Hazari, L. K. Das, A. Bauza, A. Frontera and A. Ghosh, *Dalton Trans.*, 2014, **43**, 8007; (b) P. Seth, A. Bauza, A. Frontera and A. Ghosh, *Inorg. Chem. Commun.*, 2014, **41**, 1; (c) A. Bhattacharyya, P. K. Bhaumik, A. Bauza, P. P. Jana, A. Frontera, M. G. B. Drew and S. Chattopadhyay, *RSC Adv.*, 2014, **4**, 58643; (d) H. S. Jena, *RSC Adv.*, 2014, **4**, 3028; (e) H. S. Jena, *Inorg. Chim. Acta*, 2014, **410**, 156; (f) S. Naiya, M. G. B. Drew, C. Estarellas, A. Frontera and A. Ghosh, *Inorg. Chim. Acta*, 2010, **363**, 3904; (g) C. Murcia-García, A. Bauzá, G. Schnakenburg, A. Frontera and R. Streubel, *CrystEngComm*, 2015, **17**, 1769.
- 16 K. P. Sarma and R. K. Poddar, *Transition Met. Chem.*, 1984, **9**, 135–138.
- 17 (a) G. M. Sheldrick, *Acta Crystallogr., Sect. A: Fundam. Crystallogr.*, 2008, **64**, 112–122; (b) G. M. Sheldrick, *SHELXS-97 and SHELXL-97: Program for Structure Solution*, University of Göttingen, Institute für Anorganische Chemieder Universität, Göttingen, Germany, 1997.
- 18 G. M. Sheldrick, *SADABS: Software for Empirical Absorption Correction*, University of Göttingen, Institute für Anorganische Chemieder Universität, Göttingen, Germany, 1999–2003.
- 19 R. Ahlrichs, M. Bär, M. Häser, H. Horn and C. Kölmel, *Chem. Phys. Lett.*, 1989, **162**, 165–169.
- 20 (a) A. Bauzá, A. Terrón, M. Barceló-Oliver, A. García-Raso and A. Frontera, *Inorg. Chim. Acta*, 2016, DOI: 10.1016/j.ica.2015.04.028; (b) D. Sadhukhan, M. Maiti, G. Pilet, A. Bauzá, A. Frontera and S. Mitra, *Eur. J. Inorg. Chem.*, 2015, **11**, 1958–1972; (c) M. Mirzaei, H. Eshtiagh-Hosseini, Z. Bolouri, Z. Rahmati, A. Esmailzadeh, A. Hassanpoor, A. Bauza, P. Ballester, M. Barceló-Oliver, J. T. Mague, B. Notash and A. Frontera, *Cryst. Growth Des.*, 2015, **15**, 1351–1361; (d) P. Chakraborty, S. Purkait, S. Mondal, A. Bauzá, A. Frontera, C. Massera and D. Das, *CrystEngComm*, 2015, **17**, 4680–4690.
- 21 S. F. Boys and F. Bernardi, *Mol. Phys.*, 1970, **19**, 553–566.
- 22 A. E. Reed, L. A. Curtiss and F. Weinhold, *Chem. Rev.*, 1988, **88**, 899–926.
- 23 T. A. Keith, *TK Gristmill Software, AIMAll (Version 13.05.06)*, Overland Park, KS, USA, 2013.
- 24 (a) P. Bhowmik, S. Jana, P. P. Jana, K. Harms and S. Chattopadhyay, *Inorg. Chim. Acta*, 2012, **390**, 53–60; (b) S. Bhattacharya and S. Mohanta, *Inorg. Chim. Acta*, 2015, **432**, 169–175.
- 25 (a) D. Cremer and J. A. Pople, *J. Am. Chem. Soc.*, 1975, **97**, 1354–1358; (b) D. Cremer, *Acta Crystallogr., Sect. B: Struct. Sci.*, 1984, **40**, 498–500; (c) J. C. A. Boeyens, *J. Cryst. Mol. Struct.*, 1978, **8**, 317–320.
- 26 (a) J. Echeverría, G. Aullón, D. Danovich, S. Shaik and S. Alvarez, *Nat. Chem.*, 2011, **3**, 323–330; (b) D. Danovich, S. Shaik, F. Neese, J. Echeverría, G. Aullón and S. Alvarez, *J. Chem. Theory Comput.*, 2013, **9**, 1977–1991.
- 27 A. Hazari, L. K. Das, A. Bauzá, A. Frontera and A. Ghosh, *Dalton Trans.*, 2014, **43**, 8007–8015.
- 28 F. Weinhold, *J. Comput. Chem.*, 2012, **33**, 2363–2379.
- 29 A. Bhattacharyya, S. Roy, J. Chakraborty and S. Chattopadhyay, *Polyhedron*, 2016, **112**, 109–117.
- 30 (a) S. Roy and S. Chattopadhyay, *Inorg. Chim. Acta*, 2015, **433**, 72–77; (b) S. Thakurta, R. J. Butcher, C. J. Gomez-Garcia, E. Garribba and S. Mitra, *Inorg. Chim. Acta*, 2010, **363**, 3981–3986.
- 31 (a) B. Machura, I. Nawrop, R. Kruszynski and M. Dulski, *Polyhedron*, 2013, **32**, 272–284; (b) A. Bhattacharyya, P. K. Bhaumik, M. Das, A. Bauzá, P. P. Jana, K. Harms, A. Frontera and S. Chattopadhyay, *Polyhedron*, 2015, **34**, 257–269; (c) S. Chattopadhyay, M. G. B. Drew and A. Ghosh, *Polyhedron*, 2007, **26**, 3513–3522.
- 32 A. Bhattacharyya, P. K. Bhaumik, P. P. Jana and S. Chattopadhyay, *Polyhedron*, 2014, **33**, 40–45.

