



Cite this: *CrystEngComm*, 2016, 18, 7070

Intramolecular H \cdots S interactions in metal di-(isopropyl)dithiocarbamate complexes†

Alexander Angeloski,^a James M. Hook,^b Mohan Bhadbhade,^c Anthony T. Baker^d and Andrew M. McDonagh^{*a}

Networks of C–H \cdots S interactions have been discovered within the molecular structure of sodium di-(isopropyl)dithiocarbamate pentahydrate with the formula Na(C₇H₁₄NS₂) \cdot 5H₂O, revealed by single crystal X-ray diffraction. These interactions have also been investigated by *ab initio* and Hirshfeld surface analyses which show that the electron density is not symmetrical about the molecule. NMR spectroscopy in solution and solid the state showed temperature dependent restricted rotation of the isopropyl groups, which is attributed to the intramolecular C–H \cdots S interactions. The ubiquitous nature of C–H \cdots S intramolecular interactions in this class of compound is evident in the structures of other di-(isopropyl)dithiocarbamate complexes deposited in the CSD. In general, the restricted rotation in di-(isopropyl)dithiocarbamate complexes can be directly attributed to intramolecular C–H \cdots S interactions, which subsequently influence the geometry in association with steric repulsion factors.

Received 1st July 2016,
Accepted 16th August 2016

DOI: 10.1039/c6ce01475e

www.rsc.org/crystengcomm

1. Introduction

Noncovalent interactions can exert significant influence on the geometry of molecules and their associated crystal structures.¹ Hydrogen bonding is an important noncovalent interaction where the focus has traditionally been on strong acceptors such as oxygen and nitrogen.^{2–7} In contrast, sulfur is considered a weak acceptor, and the influence of H \cdots S hydrogen bonds on molecular structure has received considerably less attention.^{8–11} Such interactions can influence side chain geometries and secondary structure in biological systems (such as those involving the amino acid cysteine).^{12–16} Thus, an understanding of the nature of H \cdots S interactions is of fundamental importance. Here, we examine a low molecular weight compound to illustrate the influence of H \cdots S interactions upon both solid and solution phase geometries. We demonstrate that the H \cdots S interaction has a profound effect in both regimes that effectively breaks the symmetry of the

molecule and creates a restricted rotation about several of the covalent bonds within the molecule.

Dithiocarbamate complexes have been studied extensively for their ability to coordinate a range of transition and main block elements, and for their interesting and useful properties.^{17,18} In particular, there is a significant amount of data pertaining to di-(isopropyl)dithiocarbamate (dipdte) compounds. The structure of the dipdte ligand is such that the C₂NCS₂ atoms lie within a plane. It has been known for some time that there is a disruption to the symmetry of dipdte complexes whereby the methine hydrogens are oriented in different directions relative to the C₂NCS₂ plane (Fig. 1). This orientation results in the inequivalence of the two isopropyl

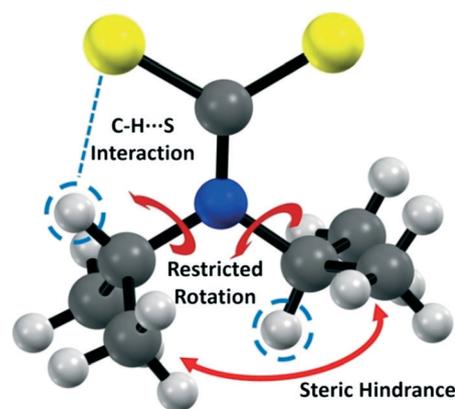


Fig. 1 Structure of the dipdte anion showing the important interactions.

^a School of Mathematical and Physical Sciences, University of Technology Sydney, Ultimo, 2007, Australia. E-mail: andrew.mcdonagh@uts.edu.au

^b NMR Facility, Mark Wainwright Analytical Centre, The University of New South Wales, Sydney, 2052, Australia

^c School of Chemistry, The University of New South Wales, Sydney, 2052, Australia

^d College of Science, Health and Engineering, La Trobe University, Melbourne, 3086, Australia

† Electronic supplementary information (ESI) available: Kinetic data, data obtained from CSD database survey of metal dipdte complexes, fingerprint plots for the Hirshfeld surface of Na(dipdte) \cdot 5H₂O, and relevant NMR spectra. CCDC 1488005. For ESI and crystallographic data in CIF or other electronic format see DOI: 10.1039/c6ce01475e



groups, which has been ascribed to a steric interaction hindering rotation about the N–C_{isopropyl} bonds based on NMR experiments in solution.^{19–23} The NMR data reveal thermodynamic parameters associated with this restricted rotation but in isolation, these studies do not elucidate the origin of the inequivalence of the two isopropyl groups.

In this work, we have redetermined²⁴ the crystal structure of sodium di-(isopropyl)dithiocarbamate pentahydrate, Na(dipdte)·5H₂O, with greater accuracy to allow a detailed examination of intramolecular C–H···S interactions. The compound was also analysed using variable temperature solution state NMR spectroscopy, which shows features associated with restricted rotation about the N–C_{isopropyl} bonds. Using a combination of theoretical calculations and experimental observations, we elucidate the origin of the inequivalence of the two isopropyl groups and restricted rotation. Furthermore, we show that these C–H···S interactions are ubiquitous throughout metal dipdte complexes for which structural determinations have been deposited.

2. Experimental

Reagents and instruments

Chemicals and solvents used in synthetic procedures were analytical grade and purchased from Sigma Aldrich and used as received. Millipore water (18.4 MΩ cm⁻¹) was used in synthetic procedures. A high resolution mass spectrum was acquired using an Agilent 6510 Q-TOF with a mobile phase of 70% acetonitrile, 30% water.

Synthesis

Sodium di-(isopropyl)dithiocarbamate was prepared as described by Angeloski *et al.*²⁵ An aqueous solution of sodium hydroxide (10.00 g in 40 mL) was cooled with stirring to 5 °C. Diisopropylamine (25 mL) was added followed by diethyl ether (75 mL). The solution was maintained between 5–10 °C whilst carbon disulfide (25 mL) was added dropwise. A precipitate formed upon addition and the resultant mixture was stirred for 20 minutes. The crude material was collected by vacuum filtration, and the filter cake purified by recrystallization using layer diffusion of ether into a methanolic solution of the crude material. The colourless crystals were collected by vacuum filtration, washed with warm diethyl ether and dried *in vacuo* to produce 15.34 g of sodium di-(isopropyl)dithiocarbamate (31%). HRMS (M + H)⁺ for NaNS₂C₇H₁₄ calculated: 200.0538; found: 200.0549. ¹H NMR (600.1 MHz, CD₃CN, 258 K): δ 6.21 (sept, *J* = 6.8 Hz, 1H, H2), 3.79 (sept, *J* = 6.9 Hz, 1H, H5), 1.63 (d, *J* = 6.8 Hz, 6H, C6H₃, C7H₃), 1.06 (d, *J* = 6.9 Hz, 6H, C3H₃, C4H₃). ¹³C NMR (150.9 MHz, CD₃CN, 258 K): δ 212.4 (C1), 56.1 (C2), 50.0 (C5), 20.9 (C6, C7), 19.6 (C3, C4).

NMR spectroscopy and kinetic analysis

Variable temperature proton (¹H), carbon (¹³C), 2D exchange spectroscopy (EXSY, with mixing time 100 ms), heteronuclear

single quantum correlation (HSQC) and heteronuclear multiple bond correlation (HMBC) nuclear magnetic spectroscopy of solutions was performed using a Bruker Avance III NMR spectrometer fitted with a BBFO Plus solution state probe. The frequency was 600.1 MHz for ¹H, 150.9 MHz for ¹³C and 60.8 MHz for ¹⁵N experiments. A Bruker BVT3000 VT unit was used in conjunction with a BCU-Xtreme cooler to accurately adjust the sample temperature between 293 and 258 K with deuterated acetonitrile as the solvent. Instrument broadening was accounted for using acetonitrile residual proton resonances. Using this data, experimental rate constants were calculated at each temperature using the signal FWHM for the methyl peak at 1.63 ppm. Arrhenius and Eyring activation parameters were obtained using generalized least squares linear regressions of log *k* vs. 1/*T* and log(*k*/*T*) vs. 1/*T* respectively. A solid state ¹H NMR spectrum was acquired using a Bruker Avance III 700 MHz solid state NMR spectrometer. Samples were loaded into a 1.3 mm zirconia rotor and a MAS rate of 60 kHz was adopted using a Bruker MAS 2 unit.

Crystallographic analysis

Crystals of Na(dipdte)·5H₂O suitable for analysis were prepared by layer diffusion of ether into a methanolic solution. A suitable crystal was selected under a polarising microscope (Leica M165Z), mounted on a MicroMount (MiTiGen) consisting of a thin polymer tip with a wicking aperture. The X-ray diffraction measurements were carried out on a Bruker Kappa-II CCD diffractometer at 150 K by using IμS Incoatec Microfocus Source with Mo-Kα radiation ($\lambda = 0.710723 \text{ \AA}$). The structure was solved using charge flipping and the full matrix least squares refinement was performed using ShelXL²⁶ in Olex2.²⁷ Heavy atoms were refined isotropically until *R*-factor convergence, and then an anisotropic model was applied. Hydrogen atoms were located using a difference Fourier plot, and restrained to neutron diffraction distances where required for water molecules.

CrystalExplorer²⁸ was used to generate Hirshfeld surfaces^{29–31} representing *d*_{norm} and electron deformation density. The latter surface was calculated using TONTO³² which is integrated into CrystalExplorer, and the experimental geometry was used as the input. The electron deformation density was mapped on the Hirshfeld surface using the 6-311G(*d,p*) basis set with the Density Functional Theory.

Crystal structure retrieval

Previous determinations of metal dipdte crystal structures were retrieved from the Cambridge Structural Database (CSD)³³ with the following specifications: *R* ≤ 5%, no disorder, no errors, no powder structures and complete 3D coordinates. This produced a total of 32 crystal structures, which were sorted by hand to remove duplicates and non-relevant structures. A final number of 28 crystal structures were retrieved. Lists of the crystal structure CSD reference codes are available in the ESI.†



Evaluation of intramolecular contacts

Atoms from the retrieved structures were relabelled to be in accordance with the labelling scheme in Fig. 2. The presence of an intramolecular C–H...S interaction was preliminarily assessed by calculation of the S1...H2 distance. Quantitative measurements were performed using the heavy atom structural parameters to avoid uncertainty due to hydrogen atom treatment. The structures were sorted by hand to retrieve the following parameters: C1–N1–C2 and C1–N1–C5 bond angles. The C1–N1, C2–S1 and C5–S2 (Fig. 2) distances were also extracted. Bond angles were kept to two decimal places, and bond lengths to three. All statistical analyses were conducted using the statistical analysis software, R 3.2.5.³⁴

3. Results and discussion

X-Ray structure determination

Relevant crystal data, selected bond lengths and angles are given in Tables 1 and 2 (see ESI† for complete data). These values are in good agreement with the previously reported structure (CSD-Refcode BUNPIJ), with bond lengths falling in the ranges expected on the analysis of the literature.^{24,36} The asymmetric unit, Fig. 2, contains one molecular anion together with one sodium cation (Na1) co-ordinated to five water molecules (O1–O5) (the sixth octahedral site is occupied by O3 on the symmetry related water molecule O3 (1 – x, 1 – y, 1 – z)). The co-ordination sphere of sodium is distorted octahedral with deviations from ideal octahedral geometry of less than 5°. The Na–O distances range from 2.3859(15) Å to 2.4492(14) Å. There is no direct S2–Na1 bond; Na1 is located more than 4 Å from S2. Within the dipdte anion, the maximum deviations from the least-squares plane through C2–C5–S2–S1 (with a r.m.s. deviation of 0.008 Å) are 0.009 Å for C2 and C5, and 0.008 Å for S1 and S2. The S2–C1 and S1–C1 bond lengths are inequivalent with S1–C1 [1.7484(16) Å] elongated compared to S2–C1 [1.7145(17) Å]. The N1–C2 and N1–C5 bond lengths are equal within experimental error and all

Table 1 Crystal data and structure refinement parameters for Na(dipdte)·5H₂O

Chemical formula	2(C ₇ H ₁₄ NS ₂)·H ₂ O·Na ₂ O ₁₀
<i>M_r</i>	578.76
Crystal system, space group	Triclinic, <i>P</i> 1̄
Temperature (K)	150
<i>a</i> , <i>b</i> , <i>c</i> (Å)	5.9472(11), 7.7189(16), 17.425(3)
<i>α</i> , <i>β</i> , <i>γ</i> (°)	92.183(8), 95.095(8), 106.851(8)
<i>V</i> (Å ³)	760.8(3)
<i>Z</i>	1
Radiation type	Mo Kα
<i>μ</i> (mm ⁻¹)	0.38
Crystal size (mm)	0.41 × 0.09 × 0.04
Absorption correction	Multi-scan <i>SADABS2014/5</i> (ref. 35) was used for absorption correction. <i>wR₂(int)</i> was 0.1663 before and 0.0594 after correction. The ratio of minimum to maximum transmission is 0.9019. The <i>λ</i> /2 correction factor is 0.00150
Diffractionmeter	Bruker <i>APEX-II</i> CCD
<i>T_{min}</i> , <i>T_{max}</i>	0.672, 0.746
No. of measured, independent and observed [<i>I</i> > 2σ(<i>I</i>)] reflections	18 709, 3304, 2666
<i>R_{int}</i>	0.060
(sin <i>θ</i> / <i>λ</i>) _{max} (Å ⁻¹)	0.639
<i>R</i> [<i>F</i> ² > 2σ(<i>F</i> ²)], <i>wR</i> (<i>F</i> ²), <i>S</i>	0.032, 0.070, 1.04
No. of reflections	3304
No. of parameters	241
No. of restraints	11
H-Atom treatment	All H-atom parameters refined
Δ _{max} , Δ _{min} (e Å ⁻³)	0.30, -0.25

bond lengths are similar to those in other M(dipdte) structures, where M = Pb, Hg, Ni, In, Co, Be, Cu and Au.^{37–44}

From the molecular assembly diagram, Fig. 3, a layered supramolecular motif is evident parallel to the crystallographic *b* direction (see ESI† for images showing packing

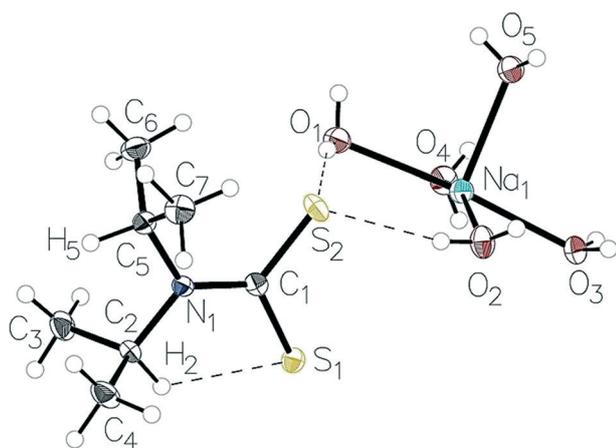


Fig. 2 The structure of the Na(dipdte)·5H₂O asymmetric unit showing the atom-labelling scheme and thermal displacement ellipsoids at 50% probability.

Table 2 Selected bond lengths (Å) and angles (°) for Na(dipdte)·5H₂O

S1–C1	1.7484(16)	Na1–O4	2.3891(13)
S2–C1	1.7145(17)	Na1–O1	2.4492(14)
N1–C2	1.492(2)	Na1–O2	2.4207(13)
N1–C1	1.345(2)	Na1–O5	2.3859(15)
C2–C3	1.524(2)	Na1–O3 ⁱ	2.4278(14)
C5–C6	1.522(2)	Na1–O3	2.3879(14)
C2–N1–C5	113.38(12)	O4–Na1–O1	92.33(5)
C1–N1–C2	122.01(13)	O4–Na1–O2	162.67(5)
C1–N1–C5	124.59(14)	O4–Na1–O3 ⁱ	84.60(5)
N1–C2–C3	111.51(13)	O2–Na1–O1	95.54(5)
N1–C2–C4	111.23(13)	O2–Na1–O3 ⁱ	80.40(5)
C4–C2–C3	112.74(15)	O5–Na1–O4	96.85(5)
S2–C1–S1	118.11(9)	O5–Na1–O1	88.03(5)
N1–C1–S1	120.25(12)	O5–Na1–O2	98.83(5)
N1–C1–S2	121.63(12)	O5–Na1–O3	97.38(5)
N1–C5–C7	113.60(13)	O5–Na1–O3 ⁱ	175.04(5)
N1–C5–C6	113.02(13)	O3–Na1–O4	85.89(5)
C6–C5–C7	113.36(15)	O3–Na1–O1	174.47(5)
O3–Na1–O2	84.83(5)	O3 ⁱ –Na1–O1	87.17(5)
Na1–O3–Na1 ⁱ	92.55(5)	O3–Na1–O3 ⁱ	87.45(5)

Symmetry code: (i) $-x + 1, -y + 1, -z + 1$.



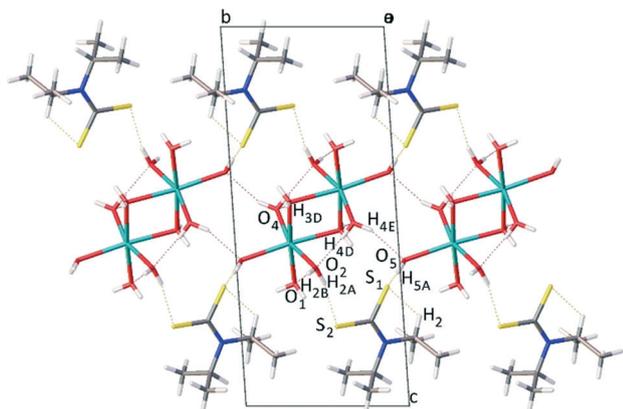


Fig. 3 Molecular packing diagram as viewed along the *a* axis, showing layered morphology of Na(dipdct)·5H₂O.

diagrams viewed along *b* and *c* directions). The layered structure is stabilised by van der Waals interactions between the alkyl groups on the anion.

Sodium ions are positioned between layers of symmetrically equivalent ligand anions, forming a two dimensional Na–Na array oriented parallel to the crystallographic *b* direction. The sodium ions are separated by a distance of 3.4804(12) Å, significantly longer than the sum of ionic radii of 2.32 Å. Alternating sodium ions are linked by bridging water molecules at O3, forming a four membered Na1–O3–Na1–O3 ring with vertices of 92.55(5)° for Na1–O3–Na1 and 87.45(5)° for O3–Na1–O3. That is, dinuclear entities [Na₂(OH₂)₁₀] exist in which the distorted NaO₆ octahedra share an edge. The array of Na cations is stabilised by a network of hydrogen bonding⁴⁵ between alternate and neighbouring water molecules where distances and angles are in agreement with literature values for O–H···O hydrogen bonds.^{46–48} The ligand anions are linked by O–H···S hydrogen bonds^{2,49} to the water molecules associated with the array of sodium cations. These hydrogen bonding contacts are summarised in Table 3.

Contacts involving intermolecular H···S, H···O and H···H interactions were examined using Hirshfeld surface analysis.^{29–31,50,51} The majority of interactions within the structure are dominated by van der Waals H···H interactions

Table 3 Selected hydrogen bond parameters

<i>D</i> –H··· <i>A</i>	<i>D</i> –H (Å)	H··· <i>A</i> (Å)	<i>D</i> ··· <i>A</i> (Å)	<i>D</i> –H··· <i>A</i> (°)
C2–H2···S1	0.998(16)	2.392(16)	3.0264(17)	120.8(12)
C7–H7A···S2	0.921(19)	2.714(17)	3.270(2)	119.7(13)
C6–H6A···S2	0.938(19)	2.741(17)	3.280(2)	117.4(13)
O2–H2A···S2	0.85(2)	2.41(2)	3.2336(15)	165(2)
O5–H5A···S1 ⁱ	0.83(2)	2.53(2)	3.2831(14)	151(2)
O3–H3D···O4 ⁱⁱ	0.83(2)	2.02(2)	2.8407(18)	171(2)
O2–H2B···O1 ⁱⁱ	0.84(2)	1.97(2)	2.8123(18)	175(2)
O4–H4D···O2 ⁱⁱⁱ	0.83(2)	1.95(2)	2.7796(19)	176(2)
O4–H4E···O5 ^{iv}	0.83(2)	2.08(2)	2.8751(19)	160(2)

Symmetry codes: (i) *x*, *y* + 1, *z*; (ii) *x* + 1, *y*, *z*; (iii) *−x* + 1, *−y* + 1, *−z* + 1; (iv) *−x* + 1, *−y* + 2, *−z* + 1.

(~65%) followed by O–H···S (~17%) and O–H···O (~13%) interactions. The intermolecular contacts are highlighted in the map of d_{norm} on the Hirshfeld surface, Fig. 4. The dark red regions are due to hydrogen bonding whilst the blue and white regions reflect H···H interactions. C–H···S interactions are also evident.

Of significance to the current work are the intramolecular non-bonding interactions involving sulfur. The S1···H2 intramolecular interaction has a distance of ~2.4 Å (Table 3), which as we show here, exerts influence throughout the entire molecular structure. The C1–N1–C2 angle is 2.58 (13)° smaller than the C1–N1–C5 angle and the C2–S1 distance is 0.034 Å shorter than the C5–S2 distance (Table 4). The S1···H2 interaction also results in a pair of interactions between S2 and the methyl hydrogens attached to C6 and C7 of ~2.7 Å (Table 3). That is, the S1···H2 interaction creates an inequivalence of the two isopropyl groups within the dipdct anion. The intramolecular C–H···S dihedral angles and lengths are similar to those involving cysteine (117.4° and 2.51 Å) and methionine (119.0° and 2.74 Å) residues interacting within proteins.¹²

Electrostatic deformation density and topological features of the intramolecular C–S···H interaction.

A 3D electrostatic deformation density map (Fig. 5) shows the electron density about the dipdct anion. The lone pair electron density (LPED) about S1 and S2 is of particular interest as this may influence hydrogen bond directionality and the capacity of S1 and S2 to form multiple H bonds.² Our calculations show that the S atom LPED adopts a toroidal geometry, in agreement with previous analyses of sulfur-containing compounds.² Upon closer examination (Fig. 6), the LPED about S1 and S2 is somewhat distorted as a result of S···H interactions. In addition to intermolecular interactions with H₂O, electron density is directed from S1 towards H2 while S2 directs electron density towards H6 and H7. A region of charge depletion (shown in red) about H2 is oriented toward the LPED about S1. This attractive behaviour between

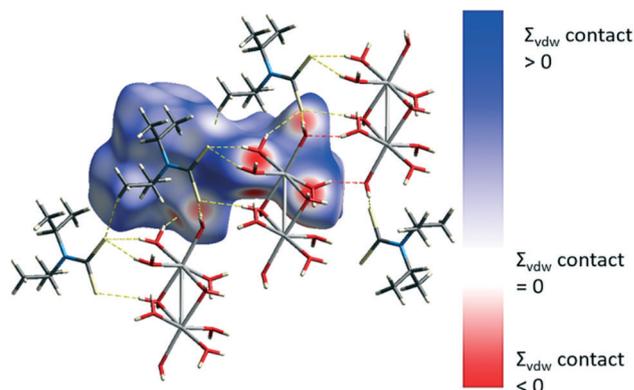


Fig. 4 Hirshfeld surface and O–H···S hydrogen bonds (yellow) and O–H···O hydrogen bonds (red).



Table 4 Summary of CSD structural parameters

Parameter	Mean \pm standard error	Current study
C2–S1 distance	3.024(6) Å	3.0264(17) Å
C5–S2 distance	3.147(8) Å	3.1014(18) Å
S2–C1–S1 angle	111.81(71)°	118.11(9)°
C1–N1 distance	1.321(2) Å	1.345(2) Å
C1–N1–C5 angle	123.78(11)°	124.59(13)°
C1–N1–C2 angle	120.11(22)°	122.01(13)°

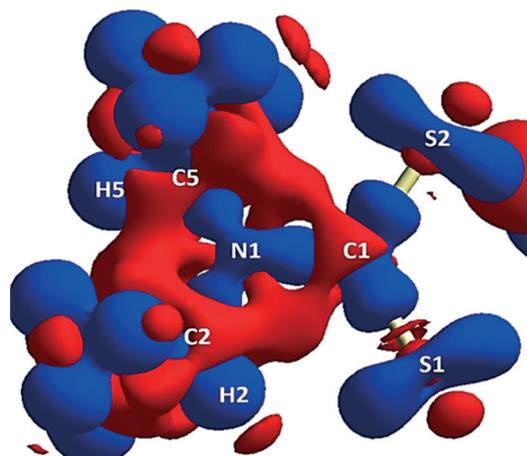


Fig. 5 Theoretical 3D electrostatic densities of the Na(dipdte)·5H₂O anion at the B3LYP 6-311G(d,p) level. Areas of lower density are shown in red, and areas of higher density are shown in blue.

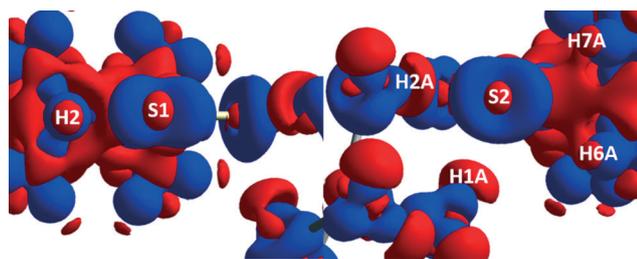


Fig. 6 3D Static deformation densities of the sulfur tori (viewed down the C–S bond).

S1 and H2 is the origin of the unsymmetrical geometry of the isopropyl groups about N1.

NMR Spectroscopy

The crystallographic intramolecular C–H \cdots S interactions yield two inequivalent isopropyl groups which are also manifested in solution, evidenced by two sets of signals in solution state NMR spectra. ¹H and ¹³C NMR spectra of Na(dipdte)·5H₂O recorded using deuterated acetonitrile at room temperature revealed broad proton and carbon resonances associated with the methyl and methine groups. Two broad signals for the methyl protons and two broad signals for the methine protons were observed, consistent with two inequivalent environments (ESI[†]). That is, H2 is maintained in synchronous gear-

like arrangement relative to H5 such that the steric repulsion between the associated methyl groups is minimised, Fig. 1. A DOSY experiment confirmed that the signals arose from only a single molecular species (ESI[†]).

The methine protons, H2 and H5, exhibit chemical shifts of 6.2 and 3.8 ppm, respectively. We attribute the large difference in chemical shifts to the influence of the C–H \cdots S interaction upon the local electron density about H2, whereby H2 is significantly more deshielded than H5. Similarly, the signal assigned to the methyl protons associated with C6 and C7 appears at 1.6 ppm while the signal assigned to protons on C3 and C4 appears at 1.1 ppm. Thus, while the influence of the C–H \cdots S interaction is still apparent, the difference in chemical shifts between the methyl protons on the two inequivalent isopropyl groups is less than that between H2 and H5 due to the greater distance between the methyl protons and sulfur. In the ¹³C NMR spectrum recorded at 293 K, two sets of signals are also observed, which are significantly broadened. 2D-EXSY spectra recorded at 293 K contain H2/H5 cross peaks with intensity equal to the source peaks indicating a chemical exchange process at room temperature (ESI[†]).

Similarly, the high speed MAS solid state ¹H NMR spectrum (ESI[†]) contains methyl resonances in the range of 0–2 ppm and a signal assigned to the methine C–H \cdots S proton H2 at 6.1 ppm. A signal at \sim 3.7 ppm is assigned to the methine proton H5, which is partially obscured by a broad peak associated with water molecules.

Variable temperature ¹H NMR (Fig. 7) data also provide kinetic insights into the rotation about the C2–N1 and C5–N1 bonds. A 2D-EXSY spectrum recorded at 258 K exhibits H2/H5 cross peaks of lower intensity than those recorded at 293 K, characteristic of reduced exchange between H2 and H5, and consistent with a greater restriction of rotation of the

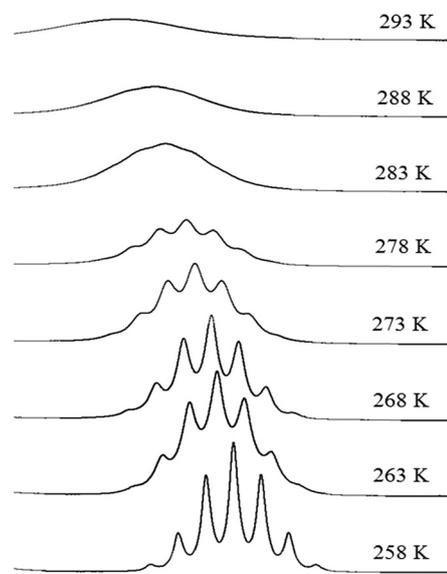


Fig. 7 Selected portion of the 600 MHz ¹H spectra for Na(dipdte)·5H₂O in CD₃CN, showing the change in H2 resonance as a function of temperature.



isopropyl groups at this temperature. Upon cooling from 293 K to 258 K, the broad signals observed in the 1D ^1H NMR spectrum resolve into their respective splitting patterns (septets for H2/H5 and doublets for the methyl protons), which is also consistent with the slowing rate of concerted rotation of the isopropyl groups. From the VT NMR data recorded using acetonitrile as solvent, a value of 63 kJ mol^{-1} for ΔG^\ddagger was calculated at 283 K and at 263 K, a value of 56 kJ mol^{-1} was obtained. While the latter value for ΔG^\ddagger agrees with data reported by others²⁰ using dichloromethane as solvent, our values for ΔH^\ddagger (27 kJ mol^{-1}), ΔS (and $-117 \text{ J mol}^{-1} \text{ K}^{-1}$) and activation energy (30 kJ mol^{-1}) are all somewhat smaller than those reported.²⁰ This indicates that the barrier to rotation may be lower in acetonitrile although quantitative comparison to the literature data²⁰ is problematic as the exact composition of the previously reported compound was uncertain.

Database survey of intramolecular C–H \cdots S interactions in metal di(isopropyl)dithiocarbamate complexes

Having examined the CH \cdots S interactions in Na(dipdte) \cdot 5H $_2$ O, we sought to find crystallographic evidence for this interaction in other dipdte complexes. A survey of the Cambridge Structural Database was undertaken. A total of 28 suitable structure determinations were selected. All structures were planar about the S $_2$ CNC $_2$ moiety and importantly, showed evidence of intramolecular S1 \cdots H2 interactions that induce differences between the two isopropyl groups. Table 4 summarises some relevant structural parameters.

The C2 to S1 distances, x (Fig. 8), were chosen for comparison in preference to H2 to S1 distances as the former are more accurate data. Values for x are tightly clustered about the mean of 3.024 Å. The shortest distance is 2.941 Å (Nd(dipdte) $_3$ phen) and the longest is 3.117 Å (Co(dipdte) $_3$). With the exception of Co(dipdte) $_3$, all values of x fall within the 2.941–3.054 Å (a range of 0.113 Å). In comparison, the C5–S2 distances (y) varied more, and ranged from 3.072 to 3.219 Å (a range of 0.147 Å) as shown in Fig. 9. The mean x distance for the 28 structures is significantly shorter (by $0.123 \pm 0.01 \text{ Å}$) than that of y ($t = -12.233$, $df = 49.412$, $p < 2.2 \times 10^{-16}$).

Unsurprisingly, the S1–C1–S2 angles (ψ) are dependent on the coordinated metal²⁴ and span the range 106.47 to 117.31°. The angle C1–N1–C2 (σ) ranges from 117.0 to 122.3° while the angle C1–N1–C5 (φ) has a range of 122.1 to 124.9°

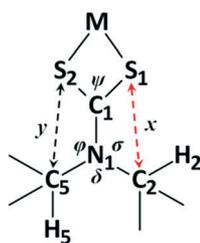


Fig. 8 Depiction of the important distances and angles discussed in the text.

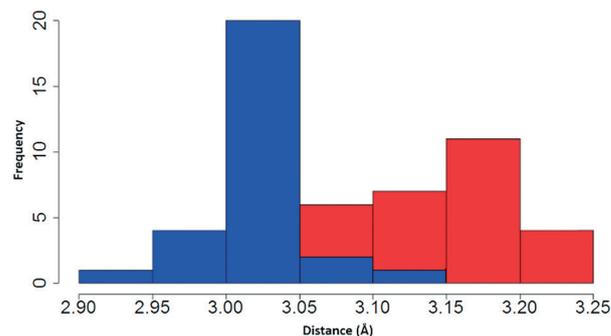


Fig. 9 Stacked histogram showing distribution of distances x (blue) and y (red).

and for any structure, $\sigma < \varphi$ ($t = -5.058$, $df = 40.716$, $p < 2.2 \times 10^{-16}$). As planarity is maintained across the S $_2$ CNC $_2$ moiety for all structures, $\varphi + \delta + \sigma = 360^\circ$ and $\delta > 114^\circ$ in all cases.

Our interpretation of these data is as follows. The angle ψ is dependent upon the coordinated metal while the distance x is a consequence of the C–H \cdots S interaction. There were no statistically significant pairwise correlations between x and ψ , nor between x and any other structural parameter. Thus, the relationships $\sigma \neq \varphi$ and $x \neq y$ are a consequence of the C–H \cdots S interaction. The C–H \cdots S interaction restricts the rotation of the corresponding isopropyl group about the C2–N1 bond. This, in turn, influences the rotation of the isopropyl group about the C5–N1 bond (through steric effects). Previously published VT NMR-based analysis of the unusual geometry of metal dipdte complexes attributed the cause of restricted rotation to purely steric effects^{19–23} but here we have shown that the underlying cause of the restricted rotation is the intramolecular C–H \cdots S interaction, an electronic effect.

Conclusions

We have shown that C–H \cdots S intramolecular interactions are present within the molecular structure of Na(dipdte) \cdot 5H $_2$ O. Theoretical calculations constrained by the experimentally derived structural data indicate that these interactions arise from the interaction between the lone pair electron density on sulfur and regions of electron depletion about a methine hydrogen.

Inequivalent chemical environments about the methine protons produced by the C–H \cdots S intramolecular interaction were observed using NMR spectroscopy in solution and solid samples. Variable temperature solution state NMR spectroscopy was used to probe the restricted rotation of the isopropyl groups about the N–C single bonds revealing an energy barrier for this rotation of 30 kJ mol^{-1} . The Gibbs free energy of the transition state (63 kJ mol^{-1}) is in agreement with previous studies of restricted rotation in dipdte structures.

An analysis of 28 similar structures using the CSD revealed the presence of intramolecular C–H \cdots S interactions. In all of the analysed structures, the heavy atom geometries supported the presence of these interactions. In all cases, the relevant intramolecular C–S distances were shorter and less



variable where intramolecular C–H⋯S interactions were present. There are no significant correlations between the steric factors of the structure and the C–H⋯S intramolecular interaction.

Thus, the restricted rotation in metal dipdte structures is directly attributable to the intramolecular C–H⋯S interaction, which subsequently influences the geometry in association with steric repulsion factors between methyl groups. We propose that these interactions are worthy of further examination in a wider range of compounds such as those found in biological systems (proteins, peptides) where bonds are subject to restricted rotation in proximity to sulfur atoms.

Acknowledgements

We acknowledge Mr Leinad Diaz and Dr Ronald Shimmon for laboratory assistance and the Mark Wainwright Analytical Centre, UNSW, for access to single crystal XRD, solid and solution state NMR spectrometers. We also acknowledge Dr Parthapratim Munshi for assistance with CrystalExplorer software.

References

- M. Nishio, Y. Umezawa, K. Honda, S. Tsuboyama and H. Suezawa, *CrystEngComm*, 2009, **11**, 1757–1788.
- B. M. Francuski, S. B. Novaković and G. A. Bogdanović, *CrystEngComm*, 2011, **13**, 3580.
- P. Schuster, G. Zundel and C. Sandorfy, *The Hydrogen Bond—Recent Developments in Theory and Experiment*, North-Holland Pub. Co, 1976.
- G. C. Pimental and A. L. McClellan, *The Hydrogen Bond*, W. H. Freeman and Co., San Francisco, 1960.
- G. A. Jeffrey and W. Saenger, *Hydrogen Bonding in Biological Structures*, Springer-Verlag, Berlin, 1991.
- P. A. Hunt, C. R. Ashworth and R. P. Matthews, *Chem. Soc. Rev.*, 2015, **44**, 1257–1288.
- A. Gavezzotti and L. L. Presti, *Cryst. Growth Des.*, 2016, **16**, 2952–2962.
- T. Steiner, *Crystallogr. Rev.*, 2003, **9**, 177–228.
- G. A. Jeffrey, *Crystallogr. Rev.*, 2003, **9**, 135–176.
- J. Perlstein, *J. Am. Chem. Soc.*, 2001, **123**, 191–192.
- G. R. Desiraju and T. Steiner, *The Weak Hydrogen Bond: In Structural Chemistry and Biology*, Oxford University Press, 2001.
- P. Zhou, F. Tian, F. Lv and Z. Shang, *Proteins: Struct., Funct., Bioinf.*, 2009, **76**, 151–163.
- D. Pal and P. Chakrabarti, *J. Biomol. Struct. Dyn.*, 2001, **19**, 115–128.
- L. M. Gregoret, S. D. Rader, R. J. Fletterick and F. E. Cohen, *Proteins: Struct., Funct., Bioinf.*, 1991, **9**, 99–107.
- M. Iwaoka and N. Isozumi, *Molecules*, 2012, **17**, 7266.
- S. Horowitz and R. C. Trievel, *J. Biol. Chem.*, 2012, **287**, 41576–41582.
- A. A. Aly, A. B. Brown, T. M. I. Bedair and E. A. Ishak, *J. Sulfur Chem.*, 2012, **33**, 605–617.
- J. J. Steggerda, J. A. Cras and J. Willemse, *Recl. Trav. Chim. Pays-Bas*, 1981, **100**, 41–48.
- R. M. Golding, P. C. Healy, P. W. G. Newman, A. H. White and E. Sinn, *Inorg. Chem.*, 1972, **11**, 2435–2440.
- A. F. Lindmark and R. C. Fay, *Inorg. Chem.*, 1983, **22**, 2000–2006.
- Y. I. Takeda, N. Watanabe and T. Tanaka, *Spectrochim. Acta, Part A*, 1976, **32**, 1553–1556.
- S. Bhattacharya, B. K. Kanungo and S. Sahoo, *J. Coord. Chem.*, 2006, **59**, 371–378.
- A. N. Bhat, R. C. Fay, D. F. Lewis, A. F. Lindmark and S. H. Strauss, *Inorg. Chem.*, 1974, **13**, 886–892.
- I. Ymen, *Acta Crystallogr., Sect. C: Cryst. Struct. Commun.*, 1983, **39**, 874–877.
- A. Angeloski, A. T. Baker, M. Bhadbhade and A. M. McDonagh, *J. Mol. Struct.*, 2016, **1113**, 127–132.
- G. M. Sheldrick, *Acta Crystallogr., Sect. A: Found. Crystallogr.*, 2008, **64**, 112–122.
- O. V. Dolomanov, L. J. Bourhis, R. J. Gildea, J. A. K. Howard and H. Puschmann, *J. Appl. Crystallogr.*, 2009, **42**, 339–341.
- S. K. Wolff, D. J. Grimwood, J. J. McKinnon, M. J. Turner, D. Jayatilaka and M. A. Spackman, *CrystalExplorer (Version 3.1)*, University of Western Australia, 2012.
- M. A. Spackman and J. J. McKinnon, *CrystEngComm*, 2002, **4**, 378–392.
- J. J. McKinnon, M. A. Spackman and A. S. Mitchell, *Acta Crystallogr., Sect. B: Struct. Sci.*, 2004, **60**, 627–668.
- J. J. McKinnon, D. Jayatilaka and M. A. Spackman, *Chem. Commun.*, 2007, 3814–3816.
- D. Jayatilaka, D. J. Grimwood, A. Lee, A. Lemay, A. J. Russel, C. Taylor, S. K. Wolff, P. Cassam-Chenai and A. Whitton, *TONTO*, Available at: <http://Hirshfieldsurface.net/>, 2005.
- F. Allen, *Acta Crystallogr., Sect. B: Struct. Sci.*, 2002, **58**, 380–388.
- R. D. C. Team, *R: A Language and Environment for Statistical Computing*, Vienna, Austria, 2008.
- Bruker, *SADABS*, Bruker AXS Inc, Wisconsin USA, Madison, 2001.
- F. H. Allen, D. G. Watson, L. Brammer, A. G. Orpen and R. Taylor, in *International Tables for Crystallography*, John Wiley & Sons, Ltd, 2006, DOI: 10.1107/97809553602060000621.
- T. A. Rodina, A. V. Ivanov, A. V. Gerasimenko, O. V. Loseva, O. N. Antzutkin and V. I. Sergienko, *Polyhedron*, 2012, **40**, 53–64.
- M. Ito and H. Iwasaki, *Acta Crystallogr., Sect. B: Struct. Crystallogr. Cryst. Chem.*, 1979, **35**, 2720–2721.
- M. Ito and H. Iwasaki, *Acta Crystallogr., Sect. B: Struct. Crystallogr. Cryst. Chem.*, 1980, **36**, 443–444.
- P. C. Healy, J. W. Connor, B. W. Skelton and A. H. White, *Aust. J. Chem.*, 1990, **43**, 1083–1095.
- S. Bhattacharya, N. Seth Miss, V. D. Gupta, H. Nöth and M. Thomann, *Z. Naturforsch., B: J. Chem. Sci.*, 1994, **49**, 193.
- F. Jian, F. Bei, P. Zhao, X. Wang, H. Fun and K. Chinnakali, *J. Coord. Chem.*, 2002, **55**, 429–437.
- H. Nöth and D. Schlosser, *Chem. Ber.*, 1988, **121**, 1711–1713.
- F. Jian, Z. Wang, Z. Bai, X. You, H.-K. Fun, K. Chinnakali and I. A. Razak, *Polyhedron*, 1999, **18**, 3401–3406.



- 45 I. A. Baburin and V. A. Blatov, *Acta Crystallogr., Sect. B: Struct. Sci.*, 2007, **63**, 791–802.
- 46 X.-Z. Li, B. Walker and A. Michaelides, *Proc. Natl. Acad. Sci. U. S. A.*, 2011, **108**, 6369–6373.
- 47 R. Kumar, J. R. Schmidt and J. L. Skinner, *J. Chem. Phys.*, 2007, **126**, 204107.
- 48 I. D. Brown, *Acta Crystallogr., Sect. A: Cryst. Phys., Diffr., Theor. Gen. Crystallogr.*, 1976, **32**, 24–31.
- 49 E. Arunan, G. R. Desiraju, R. A. Klein, J. Sadlej, S. Scheiner, I. Alkorta, D. C. Clary, R. H. Crabtree, J. J. Dannenberg, P. Hobza, H. G. Kjaergaard, A. C. Legon, B. Mennucci and D. J. Nesbitt, *Pure Appl. Chem.*, 2011, **83**, 1637–1641.
- 50 E. N. M. Yusof, M. M. Jotani, E. R. T. Tiekink and T. B. S. A. Ravooof, *Acta Crystallogr., Sect. E: Crystallogr. Commun.*, 2016, **72**, 516–521.
- 51 C. Jelsch, K. Ejsmont and L. Huder, *IUCrJ*, 2014, **1**, 119–128.

