Structural versatility of the quasi-aromatic Möbius type zinc(II)-pseudohalide complexes – experimental and theoretical investigations†

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In this contribution we report for the first time fabrication, isolation, structural and theoretical characterization of the quasi-aromatic Möbius complexes \([\text{Zn(NCS)}_2\text{Li}]_2\) (1), \([\text{Zn}_2(\mu_1,1-\text{N}_2\text{J}_2)\text{Li}_3]\) \([\text{ZnCl}_3(\text{MeOH})]_2\), 6MeOH (2) and \([\text{Zn(NCS)}_2\text{Li}]_2[\text{Zn(NCS)}_4]\) MeOH (3), constructed from 1,2-diphenyl-1,2-bis(phenylpyridin-2-yl)methylene)hydrazono)ethane \((\text{LI})\) or benzilis(bis(phenylpyridin-2-yl)methylidene)hydrazono)ethane \((\text{LII})\), respectively, and \(\text{ZnCl}_2\) mixed with \(\text{NH}_3\text{NCS}\) or \(\text{Na}_3\text{N}\). Structures 1–3 are dictated by both the bulkiness of the organic ligand and the nature of the inorganic counter ion. As evidenced by single crystal X-ray diffraction data species 1 has a neutral discrete heteroleptic mononuclear structure, whereas, complexes 2 and 3 exhibit a salt-like structure. Each structure contains a \(\text{Zn}^{2+}\) atom chelated by one tetradentate twisted ligand \((\text{LI})\) creating the unusual Möbius type topology. Theoretical investigations based on the EDDB method allowed us to determine that it constitutes the quasi-aromatic Möbius motif where a metal only induces the \(\pi\)-delocalization solely within the ligand part: 2.44\(\text{e}^-\) in 3, 3.14\(\text{e}^-\) in 2 and 3.44\(\text{e}^-\) in 1. It is found, that the degree of quasi-aromatic \(\pi\)-delocalization in the case of zinc species is significantly weaker (by \(~50\%) than the corresponding estimations for cadmium systems – it is associated with the \(\text{Zn}--\text{N}\) bonds being more polar than the related \(\text{Cd}--\text{N}\) connections. The ETS-NOCV showed that the monomers in 1 are bonded primarily through London dispersion forces, whereas long-range electrostatic stabilization is crucial in 2 and 3. A number of non-covalent interactions are additionally identified in the lattices of 1–3.

Introduction

Helical molecules are highly favoured by nature. Such molecules are of great importance, which is also supported by the structure of deoxyribonucleic acid first discovered in 1953. On the other hand, zinc(II) \((\text{Zn}^{2+})\) ions are found in all six main classes of metalloenzymes and are essential for living organisms. Moreover, the dinuclear \(\text{Zn}^{2+}\) complex fabricated from doubly deprotonated octaethyl formylbiliverdin is the first established helical doublestranded structure, which was reported in 1976. Following this discovery, strategies towards helical structure as well as their self-assembly have been the focus of researchers. Obviously, the most powerful strategy towards metal-based helical structures is the smart predesign of parent ligands. The other strategy, which, however, is less predictable and thus less efficient, is the choice of a metal-containing precursor. The latter is much less investigated.

Some efforts have been focused on the design and preparation of helical metal complexes by applying chelating ligands with suitable donor sites. The point is that the ligand should produce a helical topology upon binding to the metal ions.
some cases, the coordination features of the cations dictate the wrapping of non-helical chelating ligands around them in such a manner that they can be twisted and eventually form helical complexes. However, synthesis of organic ligands with a helical topology is more difficult than metallo-organic compounds and there are only a few reported synthetic helical organic molecules. Researchers mainly focused on the design and construction of metal complexes with synthetic helical chelating ligands. Recently, we have also directed our attention to Schiff bases comprising two pyridyl-imine functions obtained from benzylidihydrazone. These ligands were found to be efficient for helical structures upon coordination to metal centers. Particularly our comprehensive efforts were directed to various Cd salts as complexing agents. Moreover, we were able to demonstrate for the first time that the helical motif in the obtained complexes together with the chelating metalloring correspond to a quasi-aromatic Möbius object.

Herein, we report Zn(NCS)$_2$ and Zn(N$_3$)$_2$-derived structures with 1,2-diphenyl-1,2-bis((phenyl)(pyridin-2-yl)methylene) hydrazono)ethane (L$^1$) and benzilbis(acetylpyridin-2-yi) methylidenehydrazone (L$^3$). Using thiocyanate (NCS$^-$) and azide (N$_3^-$) counterions is intriguing and of great interest since both anions are known to be ambidentate ligands, which can bind metal centers in different coordination modes. As a result were able to isolate the unique quasi-aromatic Möbius type zinc complexes [Zn(NCS)$_2$L$^1$] (1), [Zn$_2$(N$_3$)$_2$(L$^3$)$_2$] (2) and [Zn(NCS)$_2$L$^3$] (3), respectively. Notably, our numerous attempts to isolate crystals of the reaction product of Zn(N$_3$)$_2$ and Zn(NCS)$_2$ failed regardless using a great number of Zn$^{II}$ and N$_3^-$ sources. DFT experiments were additionally performed to estimate the stability and aromaticity of the obtained species.

Results and discussion

Interaction of ZnCl$_2$ mixed with NH$_4$NCS or NaN$_3$ with L$^1$ or L$^3$ in MeOH at 60 °C has allowed to isolate complexes 1–3 (Scheme 1 and ESIF). The elemental analysis data supports their compositions. Notably, the same one-pot reaction of L$^1$ or L$^3$ with Cd(NO$_3$)$_2$ in the presence of NH$_4$NCS produced a dinuclear structure [Cd$_2$(N$_3$)$_2$(NCS)$_2$(L$^3$)$_2$].4MeOH (4), where L$^3$ is formed upon hydrolysis of one of the 2-PyC(Ph) functions of L$^3$, and neutral mononuclear complex [Cd(NCS)$_2$L$^3$]$(\text{MeOH})$ (5), respectively.

The FTIR spectrum of 1 contains a characteristic intense band at 2073 cm$^{-1}$ attributed to the CN stretching of NCS$^-$ ions. The same stretching mode in the IR spectrum of 3 is shown as two clearly defined bands at 2066 and 2112 cm$^{-1}$, corresponding to two different types of the NCS$^-$ ligands (Scheme 1). These bands are in the typical region for the N-linked terminal NCS$^-$ ions. The N$_3^-$ anions in the FTIR spectrum of 2 are shown as an intense band at 2054 cm$^{-1}$ arising from the $v_{\text{asym}}$ stretching vibration, as well as a band at 1441 cm$^{-1}$ corresponding to the $v_{\text{sym}}$ stretching vibration. The C=N stretching vibration is at a lower energy by 40 cm$^{-1}$ for complexes 1 and 2 compared to the free ligand L$^1$, whilst a similar 20 cm$^{-1}$ difference is observed in this vibration between compound 3 and L$^{111}$, This firmly confirms the participation of the azomethine nitrogen atoms in chelate formation. The FTIR spectra of 2 and 3 further contain a broad band for the methanol at 3353 to 3443 cm$^{-1}$, respectively.

Complex 1 crystallizes in the monoclinic space group $P2_1/n$, while complexes 2 and 3 each crystallize in the triclinic space group $P1$. It is worthy to note, that 1 is isostructural to its Mn$^{II}$ and Co$^{II}$ analogues.

Complex 1 has a neutral discrete heteroleptic mononuclear structure, where the Zn$^{II}$ metal center is coordinated by one ligand L$^1$ via its two pyridyl-imine chelate functions as well as two N-bound NCS$^-$ anions giving rise to the ZnN$_6$ chromophore with a distorted trigonal-prismatic coordination environment around the cation (Fig. 1, Table S1 in the ESIF†), which has been proven by the SHAPE 2.1 software.

Complexes 2 and 3 each exhibit a salt-like structure (Scheme 1), as opposed to our previously studied Cd$^{II}$ based counterparts. In 2, the cationic part exhibits a doubly charged centrosymmetric dinuclear structure, were two Zn$^{II}$ centers are interlinked via two $\mu_1$-$\mu_3$N$_3^-$ anions and the coordination domain of each metal is filled by the tetracoordinated ligand L$^3$ (Fig. 2). Here a coordination geometry is best described as a distorted octahedron (Table S1 in the ESIF†). The anionic part represents a discrete mononuclear structure of the composition [ZnCl$_4$(MeOH)]$_2$ with a tetracoordinate Cl$_4$O environment around the metal atom (Fig. 2). As evidenced from the so-called distortion index $\tau_4 = 0.9513$, the coordination core of [ZnCl$_4$(MeOH)]$_2$ is almost a perfect tetrahedron. This is also supported by the SHAPE 2.1 software (Table S1 in the ESIF†).

The anionic part of 2 via one of its chlorine atoms and the methanol OH hydrogen atom is engaged in intermolecular hydrogen bonds with the lattice MeOH molecules yielding a synthon of motif $R_2^1(20)$ of the ([ZnCl$_4$(MeOH)]$_2$).6MeOH composition (Fig. 2, Table S2 in the ESIF†).

The salt like structure of 3 is built from two [Zn(NCS)$_2$L$^{111}$] cations, where the Zn$^{II}$ metal center is, similar to 1 and 2, chelated by two pyridyl-imine fragments of one parent ligand L$^3$, and further bound by one N-linked NCS$^-$ anion, exhibiting a pentacoordinated geometry (Fig. 3). The distortion index $\tau_4$ is 0.693 and 0.653 for two [Zn(NCS)$_2$L$^{111}$] cations. These values are best described as being about 31% and 35%, respectively, along the pathway of distortion from the ideal trigonal bipyramidal structure towards square pyramidal structure. The trigonal bipyramidal coordination environment is also evidenced from the SHAPE 2.1 software (Table S1 in the ESIF†).

The anionic part of 3 exhibits a doubly charged [Zn(NCS)$_2$L$^{111}$] cations, where four NCS$^-$ anions are bound via their N-atoms yielding a tetracoordinated coordination geometry around the metal atom (Fig. 3). The $\tau_4$ value of 0.9594 indicates almost a perfect tetrahedron, which has also been supported by the SHAPE 2.1 software (Table S1 in the ESIF†). Notably, the CS fragment of one of the NCS$^-$ anions is disordered over three positions with a ratio of 35%: 35%: 30% (Fig. 3).

The Zn–N bonds in 1–3, formed by four nitrogen atoms of the corresponding organic ligands, are in the range from 2.3764–23775 | 23765
2.073(3) Å to 2.2960(15) Å, maintaining that Zn−NPy < Zn−Nimine (Table 1). It is worthy to note, that the Cd(L)−NCS bonds (1.970(3)−2.0289(18) Å) in 1 and 3 are remarkably shorter than the Zn−NPy and Zn−Nimine bonds, while the Zn(L)−N3 bonds in 2 are similar to those within the corresponding organic ligand. This is obviously explained by the terminal coordination mode of the NCS− anions in contrast to the μ1,1-coordination mode of N3−. The Zn−NCS bonds within the [Zn(NCS)4]2− anion in 3 are 1.953(6)−1.983(4) Å. All the NCS− and N3− ligands are almost linear (Table 1). The Zn⋯Zn separation within the dinuclear molecule of 2 is 3.3729(15) Å.

Organic ligands in the structures of 1−3 each produce a twisted geometry of different extent. As a result of this conformation, the N−C(Ph)−C(Ph)−N fragments adopt a torsion angle of about 65° (Table 1), which is significantly lower than in

Fig. 1 Crystal structure of 1 (H-hydrogen atoms are omitted for clarity). Color code: C = gold, N = blue, S = orange, Zn = magenta.

Fig. 2 (top) Crystal structure of the cationic part [Zn2(μ1,1−N3)2(L)2]2+ of 2. Hydrogen atoms are omitted for clarity. Color code: C = gold, N = blue, Zn = magenta. (bottom) Crystal structure of the hydrogen bonded synthon of motif R88(20) of the [ZnCl3(MeOH)]2−·6MeOH composition, constructed from the [ZnCl3(MeOH)]2− anionic part and lattice MeOH molecules of 2. Color code: H = black, C = gold, Cl = green, O = red, Zn = magenta.
Fig. 3  Crystal structure of 3 (hydrogen atoms and MeOH molecules are omitted for clarity). Color code: C = gold, N = blue, S = orange, Zn = magenta.

Table 1  Selected bond lengths (Å) and angles (°) for 1–3

| Bond angles | Complex 1 | Complex 2 | Complex 3 |
|-------------|-----------|-----------|-----------|
| Zn–N<sub>Py</sub> | 103.76(17) | 104.50(14), 104.63(15) | |
| Zn–N<sub>mine</sub> | 75.27(17), 75.95(17), 102.79(17), 156.78(18) | 76.44(12), 76.71(13), 78.20(14), 78.36(14), 119.93(12), 121.35(14), 163.40(13), 164.34(13) | |
| N<sub>mine</sub>–Zn–N<sub>mine</sub> | 81.62(17) | 86.81(12), 87.76(12) | |
| Zn–N<sub>mine</sub> | 86.22(7), 87.06(7), 87.39(6), 89.72(7) | 97.89(14), 98.68(13), 122.76(15), 124.22(15) | |
| N<sub>mine</sub>–Zn–N<sub>NCs</sub> | 88.08(6), 93.14(7), 144.23(6), 144.78(7) | 94.12(13), 94.20(13), 114.07(15), 114.42(16) | |
| Zn–N<sub>Py</sub> | 89.25(17), 92.07(17), 98.01(18), 161.65(17) | |
| N<sub>mine</sub>–Zn–N<sub>3</sub> | 89.89(17), 93.84(17), 105.15(17), 167.47(17) | |
| N<sub>mine</sub>–Zn–N<sub>NCs</sub> | 117.54(7) | 105.0(2), 105.3(2), 110.0(2), 112.00(17), 112.0(2), 112.73(17) | |
| N<sub>Py</sub>–Zn–N<sub>3</sub> | 79.26(17) | 165.7(4), 166.8(4) | |
| Zn–N<sub>Py</sub>–C(S) | 158.37(16), 158.46(18) | 149.5(11), 150.9(15), 158.3, 163.5(5), 168.6(4), 174.5(4) | |
| Zn–N<sub>Py</sub>–C(S) | — | — | |
| N<sub>Py</sub>–Zn–N<sub>Py</sub> | 121.8(4), 123.4(4) | — | |
| Zn–N<sub>Py</sub>–N<sub>Py</sub> | 178.62(19), 178.9(2) | 178.6(4), 179.2(4) | |
| Zn–N<sub>mine</sub>–Zn–N<sub>mine</sub> | 178.8(7) | — | |
| Zn–N<sub>mine</sub>–Zn–N<sub>NCs</sub> | 100.7(2) | — | |

Torsion angles<sup>a</sup>

| Torsion angles<sup>a</sup> | Complex 1 | Complex 2 | Complex 3 |
|---------------------------|-----------|-----------|-----------|
| N–C(Ph)–C(Ph)–N | −67.4(3) | −64.7(9) | 66.1(6), 67.6(6) |
| C(Ph)–N–N–C(Ph) | −87.0(2), −114.85(19) | −96.7(7), −143.5(6) | — |
| C(Me)–N–N–C(Ph) | — | — | 138.5(4), 140.0(4), 152.4(4), 152.9(4) |
| Py–Py | 64.19(10) | 51.13(3) | 56.6(2), 58.0(2) |
| Zn–N–N–C(Ph) | 81.04(18), 86.13(17) | 49.8(7), 81.7(5) | −47.0(5), −48.3(5), −71.2(3), −72.4(4) |

<sup>a</sup>Torsion angles must be compared by their magnitudes.
The crystal packing of 1–3 is described by a network of face-to-face π⋯π stacking between the aromatic rings (Table S3 in the ESI†). The structures of 1 and 2 are also dictated by C–H⋯π interactions (Table S4 in the ESI†).

For more detailed analyses of non-covalent interactions in 1–3 the charge and energy decomposition scheme ETS-NOCV37 is applied as available in the ADF program.28 We have applied BLYP-D3/TZP since they provide reliable results for non-covalent interactions.29 The X-ray models are considered.

We have determined, that the neutral monomers of [Zn(NCS)LII] in 1 are efficiently bonded to each other with the interaction energy ΔE\text{int} = −23.78 kcal mol⁻¹ (Fig. 4). The main gluing force (55% of the overall stabilization) is the dispersion term (ΔE\text{disp} = −23.20 kcal mol⁻¹) due to the presence of C–H⋯S and C–H⋯π contacts (Fig. 4). Such close contacts enforce additionally less important electrostatic (28.5%, ΔE\text{elstat} = −12.01 kcal mol⁻¹) and charge delocalisation (16.5%, ΔE\text{orb} = −6.95 kcal mol⁻¹) constituents (Fig. 4). The prevalence of the ΔE\text{disp} term is consistent with recent findings, which rediscover the importance of London dispersion forces in small and sizeable species.24,27,40–56

In 3 the [Zn(NCS)]²⁻ anion sticks very strongly (ΔE\text{int} = −193.21 kcal mol⁻¹) to two neighboring stacked [Zn(NCS)L]⁺ units primarily through electrostatic forces (75.3% of the overall stabilization) (Fig. 5). Quite notable (14.8%) is the charge delocalization term ΔE\text{orb} = −31.10 kcal mol⁻¹ mostly due to C–H⋯π contacts followed by the least important (9.9%) dispersion term ΔE\text{disp} = −20.62 kcal mol⁻¹ (Fig. 5). It is interesting to emphasize, that such ionic bonds are of crucial importance for the overall stability of 3 since the pure π⋯π stacking between the [Zn(NCS)L]⁺ units, though containing a significant portion of dispersion stabilization (ΔE\text{disp} = −31.76 kcal mol⁻¹), is found to be repulsive with ΔE\text{int} = 7.86 kcal mol⁻¹ caused predominantly by the unfavourable electrostatic constituent ΔE\text{elstat} = 29.1 kcal mol⁻¹ (Fig. 6). In 2 the electrostatically dominated stabilizing interactions occur between [ZnCl₂(MeOH)]⁻ and [Zn₂((μ₁,3-N3)L)₂(L)²⁺] (Fig S1 in the ESI†). Additionally, [ZnCl₂(MeOH)]⁻ forms primarily O–H⋯O as well as a series of ancillary C–H⋯Cl hydrogen bonds with the neighbouring methanol species. Such cooperative interactions, leading to ΔE\text{int} = −13.17 kcal mol⁻¹, are found to be determined mostly by the electrostatic factor (50%) followed by the charge delocalization (27%) and dispersion (23%) constituents (Fig S2 in the ESI†). The synthon [Zn₂((μ₁,3-N3)L)₂(L)²⁺] was found to be stable due to electrostatically dominated (ΔE\text{elstat} = −63.33 kcal mol⁻¹) dative-covalent Zn–N connections with ΔE\text{int} = −31.63 kcal mol⁻¹ constituted additionally from the significant portion of London dispersion forces (ΔE\text{disp} = −37.25 kcal mol⁻¹) (Fig S3 in the ESI†).

In order to evaluate aromaticity in 1–3, we have applied the electron density of delocalized bonds (EDDB) method, which has been proposed to visualize and quantify aromaticity and chemical resonance in a wide range of chemical species.57–60

Moreover, it has recently been shown that, in the case of organometalllics, the EDDB method provides very useful data on the role of the transition metal d-orbitals in electron delocalization,24,27,30,61 which is inaccessible by means of such popular and commonly used aromaticity descriptors as the nucleus-independent chemical shift (NICS)62 or the anisotropy of the induced current density (ACID).63

The global EDDB isocontours and the corresponding electron populations of 1–3 are collected in Fig. S4 in the ESI.† Here,

![Fig. 4](image_url) (top) ETS-NOCV outcomes scrutinizing the nature of bonding between the [Zn(NCS)L]⁺ monomers in 1. (bottom) The overall change in electron density Δρ\text{orb} with the corresponding energy ΔE\text{orb}.

![Fig. 5](image_url) (top) ETS-NOCV outcomes scrutinizing the nature of ionic interaction between [Zn(NCS)]²⁻ and two [Zn(NCS)L]⁺ in 3. (bottom) The overall change in electron density Δρ\text{orb} with the corresponding energy ΔE\text{orb}.
we focus our attention on the characteristic seven-membered quasi-aromatic motif (7-MR), encompassing the twisted 1,1'-(1,2-ethenediyl)bis-diazene (BDA) fragment and the metal atom (abbreviated as BDA–Zn). The BDA-based complexes with cadmium have recently been demonstrated to exhibit a unique type of transition-metal induced Möbius-like aromaticity in which the metal d-orbitals themselves do not contribute to the π-conjugation occurring at BDA.24,27 Since the quantitative study of aromaticity/electron delocalization in large systems is very difficult in practice, we have decided to consider the simplified BDA–Zn models adopting the exact fragments geometries from crystals of 1–3 (Fig. 7). The calculated total EDB contours and electron populations have been dissected (according to the orbital symmetry) to get the strict π-contributions to quasi-aromaticity; natural atomic charges on the metal and the two closest nitrogen atoms have been added together with the average dihedral angles and the calculated electric dipole moments (EDM). It was found, that the number of π-electrons delocalized in the quasi-aromatic rings, particularly in 1 and 2 (on average ~3.3|e|), resembles pretty much the values found for the previously studied BDA–Cd complexes, despite different configurations of the phenyl units and applying other ligands.23–27 The most twisted 7-MR in 1 (containing the most bulky substituents) is at the same time the most stabilized by quasi-aromaticity (3.44|e|, i.e. ~0.6|e| per a quasi-ring member, which is comparable to the corresponding value for pyrrole65). Interestingly, it is found for the first time, that the systematic increase of the Zn–N bond polarization when going from 1 to 3 reveals a strict correlation between EDM (i.e. indirectly the topology and the metal to BDA charge transfer) and π-electron delocalization: $R = -0.986$. In other words, the more twisted is the 7-MR, the more quasi-aromatic character is observed (Fig. 7). It demonstrates the two-folded role of bulky substituents: they are not only dispersion donors,29–56 but they also lead to amplification of the 7-MR twist (and enhanced quasi-aromaticity). Previously only the former feature has been recognized.23–27 Interestingly, the optimized BDA–Zn structure (without steric effects from the Ph units) has significantly reduced quasi-aromaticity compared to 1 and 2 (2.43|e|, i.e. ~0.4|e| per a quasi-ring member, which is comparable to the corresponding value for furan),66 but at the same time, it is
almost twice less aromatically-stabilized than its optimized BDA–Cd analogue (4.74 [e] , |e| ~ 0.8 |e| per a quasi-ring member, which is exactly between the corresponding values for pyrrole and benzene). Since both equilibrium structures have comparative average dihedral angles N–N–M–N, it is clearly the larger metal–nitrogen bond polarization (the charges $q_\text{Zn} = +1.15, q_\text{N} = -0.9$, EDM $= 1.26$ D in BDA–Zn compared to $q_\text{Zn} = +0.72, q_\text{N} = -0.7$, EDM $= 0.68$ D in BDA-Cd) that limits the $\pi$-electron delocalization in the 7-membered quasi-aromatic unit (changes in the effectiveness of $\pi$-delocalization involving the 2pz orbitals of nitrogen atoms at close proximity of the metals are well marked in the EDBD$_2$[r] isocountours, Fig. 7). Such interaction between the nature of metal–ligand bonding and quasi-aromaticity of the ligand has not been known before.23–27

**Conclusions**

In summary, we successfully isolated and characterized the quasi-aromatic Möbius type zinc complexes [Zn(NCS)$_2$L]$^+$ (1), [Zn$_2$(μ$_1$-N$_3$)$_3$(μ$_3$)N$_2$][ZnCl$_4$(MeOH)$_2$.6MeOH (2) and [Zn(NCS)$_4$L]$^{2-}$[Zn(NCS)$_4$L]$_{-1}$, MeOH (3), fabricated from 1,2-diphenyl-1,2-bis(phenyl(pyridin-2-yl)methylene)hydrazonophenol (L$^+$) or benzilbis(acetylmethylidenehydrazone) (L$^-$), respectively, and ZnCl$_2$ mixed with NH$_4$NCS or NaN$_3$. The creation of 1–3 is dictated by both the bulkiness of the organic ligand and the nature of inorganic counter ion.

Complex 1 has a neutral discrete heteroleptic mononuclear structure with the Zn$^{II}$ metal atom being chelated by one tetradentate ligand L$^+$ and two N-bound NCS$^-$ anions with the formation of a distorted trigonal-prismatic ZnN$_6$ coordination core. The [Zn(NCS)$_2$L]$^+$ monomers were found (due to the ETS-NOCV calculations) to be bonded to each other primarily through London dispersion forces exerted by the presence of bulky hydrophobic substituents. Contrary, complexes 2 and 3 exhibit a salt-like structure where the long-range electrostatic forces were found to be of prime importance additionally to more typical non-covalent interactions (O–H···O, C–H···Cl, C–H···S, C–H···π, π···π). In 2, the cation is a doubly charged centrosymmetric dinuclear structure with two Zn$^{II}$ atoms linked via two μ$_1$-N$_3$$^-$ anions. Each metal center is further linked by the tetracoordinate ligand L$^1$. The anionic part has a discrete mononuclear composition [ZnCl$_4$(MeOH)]$^-$. Notably, the anionic part of 2 together with the Lewis MeOH molecules produces a synthons of motif $K_g(20)$ stabilized mostly by O–H···O and C–H···Cl interactions. Complex 3 is composed from two [Zn(NCS)$_4$L]$^{2-}$ cations with the Zn$^{II}$ atoms each being chelated by two pyridyl-imine fragments of L$^-$ and further bound by one N-linked NCS$^-$ anion. The anionic part of 3 is a doubly charged [Zn(NCS)$_4$L]$^{2-}$ species. Long range electrostatic forces between [Zn(NCS)$_4$L]$^{2-}$ and [Zn(NCS)$_4$L]$^{2-}$ are responsible for the stability of 3 since pure $\pi$···$\pi$ stacking between the [Zn(NCS)$_4$L]$^{2-}$ units appeared to be repulsive. Finally, we have proven, by means of the EDBD$^{27–44}$ study, that the seven-membered rings in 1–3 constitute a quasi-aromatic Möbius-type motif, though the absolute magnitude of such $\pi$-delocalization is notably weaker than in the corresponding cadmium-based analogs.24,27 Bulkiness of the ligands (L) are found not only to amplify London dispersion stabilization,24,27,40–56 but also influence the magnitude of quasi-$\pi$-delocalization (of Möbius-type) through modification of the polarity of Zn–L bonding.

**Experimental**

**Materials**

All chemicals and solvents were used from commercial sources without further purification. L$^+$ and L$^-$ were synthesized according to a literature method.20

**Physical measurements**

FTIR spectra were recorded on a Bruker Tensor 27 FTIR spectrometer. Microanalyses were performed using an ElementarVario EL III analyzer.

**Synthesis**

ZnCl$_2$ (0.068 g, 0.5 mmol), NH$_4$NCS (0.076 g, 1 mmol) or NaN$_3$ (0.065 g, 1 mmol) and L$^+$ or L$^-$ (0.284 and 0.222 g, respectively; 0.5 mmol) were placed in the main arm of a branched tube, MeOH (15 mL) was carefully added to fill the arms. The tube was sealed and immersed in an oil bath at 60 °C while the branched arm was kept at ambient temperature. X-ray suitable crystals were formed during the next days in the cooler arm and were filtered off.

1 Colorless block-like crystals. Yield: 0.248 g (66%). Anal. calc. for C$_{10}$H$_{20}$N$_4$S$_2$Zn (750.22) (%): C 60.04, H 3.76 and N 14.94; found: C 60.29, H 3.83 and N 14.77.

2 Yellow block-like crystals. Yield: 0.156 g (64%). Anal. calc. for C$_{10}$H$_{20}$Cl$_4$N$_{16}$O$_4$Zn$_4$ (1951.98) (%): C 51.69, H 4.54 and N 12.92; found: C 51.56, H 4.61 and N 12.81.

3) Yellow prism-like crystals. Yield: 0.174 g (73%). Anal. calc. for C$_{10}$H$_{20}$N$_4$S$_2$Zn$_4$ (1465.74) (%): C 51.62, H 3.83 and N 17.20; found: C 51.76, H 3.48 and N 17.33.

**ETS-NOCV charge and energy decomposition method**

The Natural Orbitals for Chemical Valence (NOCV) $\varphi_i$ constitute the canonical representation for a differential density matrix $\Delta \rho$ (it is formed by subtracting the appropriate molecular fragments density matrices from a density matrix of a molecule under consideration) in which $\Delta \rho$ adopts a diagonal form. It gives rise to the corresponding eigenvalues $\varphi_i$ and the related vectors $\varphi_i$. NOCVs occur in pairs ($\varphi_i$, $\varphi_j$) related to $\{\varphi_i\}$ and they decompose overall deformation density $\Delta \rho$ into bonding components with different symmetries ($\Delta \rho_k$):

$$\Delta \rho(r) = \sum_{k=1}^{M/2} \varphi_i \left[ -\varphi_i^2 + \varphi_i^2(r) \right] = \sum_{k=1}^{M/2} \Delta \rho_k(r)$$

Usually, a few $k$ allow to recover a major shape of $\Delta \rho$. By combining NOCVs with ETS scheme in ETS-NOCV, one can obtain the related energetics, $\Delta E_{\text{EKin}}(k)$, in addition to qualitative picture emerging from $\Delta \rho_k$. ETS originally divides the total bonding energy, between fragments, $\Delta E_{\text{total}}$, into four distinct components: $\Delta E_{\text{total}} = \Delta E_{\text{EKin}} + \Delta E_{\text{Pauli}} + \Delta E_{\text{orb}} + \Delta E_{\text{dispersion}}$.
The $\Delta E_{\text{elstat}}$ is an energy of quasi-classical electrostatic interaction between fragments. The next term, $\Delta E_{\text{Pauli}}$, is responsible for repulsive Pauli interaction between occupied orbitals on the two fragments. The third component, $\Delta E_{\text{orb}}$, is stabilizing and shows formation of a chemical bond (including polarizations).

In the ETS-NOCV scheme $\Delta E_{\text{orb}}$ is expressed in terms of the eigenvalues $v_k$ and diagonal Fock energy matrix elements $F_{kk}^{TS}$ (transformed into NOCV representation) as:

$$\Delta E_{\text{orb}} = \sum_k \Delta E_{\text{orb}}(k) = \sum_{k=1}^{M/2} v_k \left[ -F_{kk}^{TS} + F_{kk}^{TS} \right]$$

Finally, $\Delta E_{\text{dispersion}}$ denotes the semiempirical Grimme dispersion correction (D3).

**Single-crystal X-ray diffraction**

The X-ray data were collected on a Bruker APEX-II CCD single crystal diffractometer using graphite-monochromated Mo-Kα radiation ($\lambda = 0.71073$ Å). The collected frames were integrated with the Saint$^{44}$ software using a narrow-frame algorithm. Data were corrected for absorption effects using the multi-scan method in SADABS.$^{46}$ The space groups were assigned using XPREP of the Bruker ShelXTL$^{66}$ package, solved with ShelXT$^{66}$ and refined with ShelXL$^{66}$ and the graphical interface ShelXle.$^{67}$ All non-hydrogen atoms were refined anisotropically. Hydrogen atoms attached to carbon were positioned geometrically and re

**Crystal data for 1.** C$_{48}$H$_{32}$N$_{8}$S$_{2}$Zn, $M_t = 750.19$ g mol$^{-1}$, $T = 296(2)$ K, monoclinic, space group $P2_1/n$, $a = 12.8146(10)$, $b = 20.5355(17)$, $c = 13.4451(11)$ Å, $\alpha = 98.225(1)^{\circ}$, $V = 3501.7(5)$ Å$^3$, $Z = 4$, $\rho = 1.423$ g cm$^{-3}$, $\mu$(Mo-Kα) = 0.863 mm$^{-1}$, reflections: 5447 collected, 5447 unique, $R_{\text{int}} = 0.034$, $R_1$(all) = 0.0363, $wR_2$(all) = 0.0588, $S = 1.026$.

**Crystal data for 2.** C$_{46}$H$_{32}$N$_{4}$S$_{2}$Zn$_{2}$, C$_{2}$H$_{8}$ClO$_{2}$Zn$_{2}$, 6(CH$_{4}$O); $M_t = 1951.9$ g mol$^{-1}$, $T = 125(2)$ K, triclinic, space group $P\bar{1}$, $a = 11.836(4)$, $b = 14.385(4)$, $c = 14.807(4)$ Å, $\alpha = 68.045(3)$, $\beta = 83.861(3)$, $\gamma = 78.264(3)$, $V = 2288.0(12)$ Å$^3$, $Z = 2$, $\rho = 1.417$ g cm$^{-3}$, $\mu$(Mo-Kα) = 1.274 mm$^{-1}$, reflections: 15639 collected, 5424 unique, $R_{\text{int}} = 0.053$, $R_1$(all) = 0.0648, $wR_2$(all) = 0.1386, $S = 1.092$.

**Crystal data for 3.** 2(C$_{29}$H$_{24}$N$_{2}$S$_{2}$Zn), C$_{2}$N$_{8}$S$_{4}$Zn, CO; $M_t = 1461.66$ g mol$^{-1}$, $T = 100(1)$ K, triclinic, space group $P\bar{1}$, $a = 13.3235(9)$, $b = 15.6906(10)$, $c = 18.6880(13)$ Å, $\alpha = 65.795(2)$, $\beta = 71.872(2)$, $\gamma = 77.010(2)^{\circ}$, $V = 3365.1(4)$ Å$^3$, $Z = 2$, $\rho = 1.443$ g cm$^{-3}$, $\mu$(Mo-Kα) = 1.302 mm$^{-1}$, reflections: 45317 collected, 15733 unique, $R_{\text{int}} = 0.060$, $R_1$(all) = 0.1102, $wR_2$(all) = 0.1712, $S = 1.024$.

**Contributions**

Mariusz P. Mitoraj has planned and partially performed (ETS-NOCV) the theoretical calculations, written the manuscript text and analyzed the entire data. Farhad Akbari Afkhami has primarily done the experimental part. Ghodrat Mahmaoudi has planned the experimental research. Ali Akbar Khandar has supported the work, whereas Atash V. Gurbanov has synthesized the compounds. Fedor I. Zubkov has also participated in the synthesis of the compounds. Rory Waterman is a crystallographer of compound 1-2, whereas Himanshu Sekhar Jena is a crystallographer of system 3. D W. Szczepanik has done the aromaticity calculations. Damir A. Safin and Maria G. Babashkina have analysed and discussed the results as well as have written the manuscript.

**Conflicts of interest**

There are no conflicts to declare.

**Acknowledgements**

We are grateful to the University of Maragheh for the financial support of this research. We also thank “RUDN University Program 5-100” for the support. This work was also partially supported by the National Science Centre, Poland (grant no. 2015/17/D/ST4/00558, D. W. S.). M. P. M. acknowledges the financial support of the Polish National Science Center within the Sonata Bis Project 2017/26/E/ST4/00104. X-ray facilities were provided by the U. S. National Science Foundation (CHE-1039436 to RW). H. S. J. thanks FWO [PEGASUS]2 Marie Skłodowska-Curie grant agreement no. 665501 for Incoming post-doctoral fellowship. D. W. S. acknowledges the European Union’s Framework Programme for Research and Innovation Horizon 2020 (2014–2020) under the Marie Skłodowska-Curie Grant Agreement No. 797335 “MulArEffect”.

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