Uncovering the action behind negative thermal expansion in MOF-5

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Materials with properties zero or low temperature dependence find widespread applications ranging from electronic devices to dental fillings. Negative thermal expansion (NTE) materials are compounds which contract with increasing temperature. This is in contrast to most materials, which exhibit positive thermal expansion (PTE). Composites of PTE and NTE materials can be engineered to possess specific expansion properties. Understanding the mechanism behind thermal expansion on an atomic level is crucial for tailoring properties.

PTE is caused by the anharmonicity of the chemical bond potential. In NTE materials, a physical mechanism - often of vibrational character - overrules the PTE. Vibrational NTE is typically observed in flexible and porous covalent structures, which can partly absorb thermal energy as vibrational motion perpendicular to the bond direction i.e. as transverse motion. Such motions are known to cause NTE in some oxides and cyanides. In these compounds oxygen atoms or cyanide groups link together the metal atoms. NTE was recently discovered in some metal-organic frameworks (MOF) including the archetype oxides and cyanides. In these compounds oxygen atoms or cyanide groups link together the metal atoms. NTE was recently discovered in some metal-organic frameworks (MOF) including the archetype.

Our initial studies of MOF-5 used single-crystal and powder X-ray diffraction [1]. These results have been backed up by powder and single-crystal neutron diffraction data covering the temperature range 4-400 K to obtain a more detailed picture of the NTE mechanism especially with respect to the thermal motion of the hydrogen atoms. Additionally, the metal center motion was selectively probed using X-ray absorption spectroscopy (XAS) to contrast time- and position-averaged diffraction data. The complementary findings from diffraction and XAS are shown in this presentation. Powder diffraction data reveal near-linear NTE over the temperature range 20-500 K. The vibrational motions in MOF-5 were obtained from multi-temperature single-crystal neutron Laue diffraction data (4-300 K) through extraction of positional and vibrational parameters. The exact same crystal was used in a complementary single-crystal X-ray diffraction experiment covering the temperature range 100-300 K. The crystal was a solvent-free single crystal packed in a sealed glass capillary under vacuum.

There is good overall agreement between the studies showing that NTE in MOF-5 is due to (1) concerted transverse motion of the benzenedicarboxylate linkers and (2) local motion of the metal centers. Analysis of the temperature dependent atomic displacement parameters indicates that a lattice contracting transverse motion of the aromatic ring perpendicular to the unit cell axis is significantly populated below 20 K, whereas local NTE motions of the carboxylate group become populated at slightly higher temperatures.


Keywords: host-guest, metal-organic framework, porosity

Structural analysis of metal organic frameworks with perovskite-like structure

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Hybrid materials that combine inorganic and organic components and that contain cavities and channels –the so-called metal organic frameworks (MOFs)– have been extensively studied in the last decade in view of their interesting potential applications, for example, in catalysis and gas storage [1]. In addition to nanoporous MOFs, dense hybrid framework structures have also been discovered and are receiving increasing attention as they can display a wide range of interesting functional properties [2].

A most interesting example is the recently discovered family of MOFs that crystallize in the perovskite architecture, and that display weak ferromagnetism [3], dielectric properties [4] and even multiferroic behavior [5].

As it is well-known, the “conventional” perovskite structure, of general formula ABX₃ (A= lanthanide, alkaline-earth cations, etc.; B= transition metal cations; X= O₂⁻, S₂⁻, halide ions, etc.) consists of a three-dimensional array of corner sharing [BX₆] octahedra with the larger A cation occupying the 12-coordinated cuboctahedral cavities of the 3D network. Such structure is the basis of many transition metal mixed oxides with very outstanding properties (high Tc superconductivity, colossal magnetoresistance, ferroelectricity, charge ordering, nonlinear optical behaviour, catalytic properties, etc.) and extensive work has allowed to establish important structure-properties relationships.

For the much less explored MOF compounds with perovskite architecture, up to now, all of them contain the formate ion HCOO⁻ in the X site, an amine cation in the A site, while different divalent cations, as Cu²⁺, Mn²⁺, Co²⁺, Ni²⁺, Zn²⁺ and Mg²⁺ can occupy the B site [6].

In this work, and by analogy with the “conventional” perovskite, we try to find and establish systematic trends in the structural characteristics and distortions displayed by these perovskite MOFs and that can be useful to understand and even predict the properties of these compounds.

For this purpose we present a thoroughly structural analysis that we have made on two series of compounds, namely: M(HCOO)₂[(CH₃)₂NH]₃(M²⁺: Mn²⁺ to Zn²⁺) and Mn(HCOO)₂[RH]₃ (where R is an amine), and that allow us to rationalize the different parameters that control the resulting structure.
Novel amidrazone derivative and its Cu(II) complex: Crystal structure and antitumor activity

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Compounds with an open chain amidrazone system, N=C=N–N, (Scheme) constitute a unique group of ligands with their propensity to react with a wide range of transition metals in their neutral or ionic forms as well as diversity of their coordination modes [1]. Depending on the quality of the metal center and experimental conditions they can form mono-, bi- and poly-nuclear species. These properties make them useful in design and synthesis of novel functional materials. However, the most extensive studies of hydrazones are related to their coordination chemistry on the quality of the metal center and experimental conditions they exhibit significant antibacterial and antitumor properties [2], [3].

X-ray diffraction analysis of 6-acetyl-cyclohex-3-ene carboxylic acid [1-pyridin-2-yl-1-(pyridin-2-yl)amino]methyl (Z)-ylidene) hydrazide, H$_2$L, (1) and its copper(II) complex [Cu$_2$L$_2$]$_2$H$_2$O (2) has been carried out in order to elucidate the influence of coordination and amide protonation state on the geometry of the ligand.

![Structural analysis](image)

**Structure:**

Analysis showed that compound (1) exists in its amid Hydrazone form in the solid state. The central amidrazone moiety has a Z configuration with respect to the hydrazone C=N double bond. The N=C=N–N–(C=O) chain, which adopts a cis,trans,cis conformation, is almost planar. All atoms of the amidrazone moiety may be regarded as sp$^2$ hybridized. Near-planarity of this unit may suggest a high level of π-electron delocalization. However, the X-ray data indicate, that there is a clear distinction between single and double bonds in this part of molecule.

The reaction of H$_2$L (1) with copper(II) acetate results in double protonation of the ligand, namely the carboxylic and amide groups. This induces considerable π-electron delocalization along the whole amidrazone system. Furthermore, the ligand configuration is found to be transferred from Z to E upon metal complexation. The elementary building units in crystal (2) are centrosymmetric binuclear species. Isomerization around the C=N bond allows the L$^+$ ions to chelate the Cu$^{2+}$ ion through its pyridine-N, amide-O and imine-N atoms. The carboxylic O atom from the adjacent, inversion-related ligand completes the square-planar donor arrangement around the metal center.

In cytotoxicity research, (2) shown a high in vitro cytotoxic properties against SW 948, CX-1 and A-431 cancer cell lines, whereas growth inhibition activity of the free ligand (1) was no significant.

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**Keywords:** amidrazone, copper complex, crystal structure

C386-C387

Molecular imprinting effect of the guest in β-cyclodextrin inclusion complexes

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Cyclodextrins (CDs) are well known, water soluble, cyclic oligosaccharides, which are used for micro encapsulation of organic molecules inside their relatively apolar cavity. JCD inclusion complexes of two Schiff bases (anils), derivatives of salicylideneaniline, changed the behaviour of the anils from thermochromic (as free crystalline compounds) to photochromic upon encapsulation [1]. Trials to grow single crystals of the βCD/anil complexes from absolute ethanol resulted in isomorphous crystals (P2$_1$, a=15.887(8), b=14.784(12), c=15.29.680 (14) Å, β=103.19(2)°, Z=2) exhibiting a novel packing, observed for the first time, accompanied by dramatic distortions of the βCD conformations. Refinement of both structures did not reveal the guest in a clearly detectable amount, the structures appearing as βCD-ethanol complexes. However, they differ also from any of the three known forms, I – III, of βCD/ethanol complexes [2]. The structures exhibit an open space at the secondary entrance of one of the host monomers, in which the hydrophilic salicylidene moiety of the guests can fit (Figure). It is proposed that the βCD/anil complexes were initially formed by inclusion of the hydrophobic half of the guests inside the host cavity and the salicylidene part extending outside the entrance of the secondary face. Subsequently, the anils were solvolysed thereby emptying the cavity and lattice. However due to the low solubility of βCD in ethanol, the formed crystals remained intact bearing the molecular imprint of the guest, whereas its place in the crystals has been taken eventually by ethanol molecules.


**Keywords:** amidrazone, copper complex, crystal structure