Coordination-induced reversible electrical conductivity variation in the MOF-74 analogue Fe$_2$(DSBDC)$^+$

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Introduction

Traditionally raising interest due to their highly porous nature, metal–organic frameworks (MOFs) offer an excellent platform for investigating and tuning electrical, magnetic, and optical properties in organic-based materials, some of which have already led to applications in lithium-ion batteries, thermoelectrics, and supercapacitors, among others. The electronic properties may be modulated either inherently, through changes in chemical composition, or through host-guest interactions. Guest molecules may cause structural variation, tune electrical conductivity, induce spin state transitions, modulate luminescence wavelength, and even participate in guest-guest interactions. These phenomena are of fundamental interest and form the basis of various applications including chemiresistive sensors, fluorescence sensors, and cooperative adsorption.

Although several strategies are effective for increasing the electrical conductivity in MOFs, among the most promising, and certainly more tractable synthetically, are those that involve post-synthetic doping. Guest molecules introduced after the formation of a MOF may tune either the charge mobility or the charge density in the skeleton of a given material, allowing in either case continuous enhancement of electrical conductivity over several orders of magnitude. Chemically, the dopant molecules may engage with the MOF through outer-sphere electron transfer (i.e. redox reactivity) or through inner-sphere reactivity. The latter involves binding of guest molecules to coordinatively unsaturated metal centers to form charge transport pathways and/or to inject charge carriers. The inner sphere mechanism is proposed to be operative, for instance, in TCNQ-infiltrated Cu$_3$(benzene-1,3,5-tricarboxylate)$_2$ (TCNQ = 7,7,8,8-tetracyanoquinodimethane), wherein TCNQ molecules coordinate to Cu sites and generate increasingly efficient charge percolation pathways and million-fold increase in the electrical conductivity of the parent MOF.

Herein, we show that simple coordination of solvent molecules such as DMF modulates the electrical conductivity of Fe$_2$(DSBDC)$^+$ by three orders of magnitude. Fe$_2$(DSBDC) is a structural analogue of the well-known MOF-74 series M$_2$(2,5-dihydroxybenzene-1,4-dicarboxylate) (M$_2$(DOBDC), M = Mg, Mn, Fe, Co, Ni, Cu, Zn) where phenoxide groups on the DOBDC ligand have been replaced by thiophenoxide. Each Fe atom in Fe$_2$(DSBDC) is coordinated by two trans thiophenoxide groups, three meridionally coordinated carboxylate groups, and one DMF molecule. The latter can be removed to yield coordinatively unsaturated Fe sites. The Fe and thiophenoxide S atoms form [-Fe-S$_2$]$_x$ chains, which are bridged by DSBDC$^-$ ligands to form a three-dimensional framework containing one-dimensional hexagonal pores (Fig. 1). Previously, we reported that the regular hexagonal pores of DMF-filled Fe$_2$(DSBDC) (Fe$_2$(DSBDC)(DMF)$_2$) distort significantly when the unbound guest DMF molecules are replaced by dichloromethane (DCM) and the material is activated 80 °C to give Fe$_2$(DSBDC)(DMF)$_2$. The distortion is reversible: soaking

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Fe₂(DSBDC)(DMF)₂ in DMF regenerates the regular hexagonal pores and original structure. We extend the structural and electrical properties study of this system here by investigating the fully desolvated material, Fe₂(DSBDC), which shows divergent behaviour from its solvated congeners.

**Results and discussion**

Fe₂(DSBDC)(DMF)₂·ₓ(DMF) was synthesized according to a previously reported procedure.²⁶ Owing to the sensitivity of this material to both water and air, all subsequent manipulations were performed in a dry, oxygen-free atmosphere. Thermogravimetric analysis (TGA) of this material revealed two consecutive weight loss events at 210–335 °C and 320–560 °C, respectively (Fig. S1†). The first weight loss step can be correlated to the loss of both bound and unbound DMF molecules, while the second likely corresponds to framework decomposition. Indeed, powder X-ray diffraction (PXRD) confirmed that heating Fe₂(DSBDC)(DMF)₂·ₓ(DMF) at 200 °C under vacuum for 2 days causes amorphization (Fig. S2†). In an attempt to activate the material without decomposition, we exchanged DMF by soaking the as-synthesized MOF in lower-boiling methanol (MeOH), to produce Fe₂(DSBDC)(MeOH)₂·ₓ(MeOH). Near-complete displacement of DMF by MeOH was confirmed by infrared (IR) spectroscopy, which revealed significantly decreased intensity of the C=O stretch at 1654 cm⁻¹ and appearance of a MeOH C–O stretching band at 1008 cm⁻¹ as well as a broad O–H stretching band at ~3300 cm⁻¹ (Fig. 2). Although Fe₂(DSBDC)(MeOH)₂·ₓ(MeOH) remains crystalline when kept in liquid MeOH (Fig. 3), it also decomposes upon desolvation (Fig. S2†), possibly because of a stronger interaction between MeOH and the framework. We therefore sought to replace methanol with a weaker interacting solvent and soaked the methanol-exchanged material in tetrahydrofuran (THF) to produce Fe₂(DSBDC)(MeOH)₂·ₓ(THF). IR spectroscopy once again confirmed near-quantitative exchange, with complete disappearance of the bands at 1008 and 3300 cm⁻¹ and rise of characteristic C–H stretching bands for THF at 2969 and 2866 cm⁻¹, which are blue-shifted from the C–H stretching bands for MeOH at 2938 and 2831 cm⁻¹. Note that direct replacement of DMF by THF is not successful, presumably because THF is too poorly coordinating and cannot displace Fe-bound DMF, thus requiring the intermediary of MeOH.

TGA of Fe₂(DSBDC)(THF)₂·ₓ(THF) confirmed that THF can be removed even below 100 °C, much lower than the decomposition temperature of approximately 270 °C (Fig. S3†). Therefore, to remove all solvent and produce completely activated MOF, Fe₂(DSBDC)(THF)₂·ₓ(THF) was heated at 170 °C under dynamic vacuum (<3 mTorr) for 36 h. This yielded a dark red powder whose elemental microanalysis revealed a

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**Fig. 1** Structural changes induced in Fe₂(DSBDC) by coordination and release of DMF, with respective electrical conductivity (σ) values. The structures of Fe₂(DSBDC)(DMF)₂·ₓ(DMF) and Fe₂(DSBDC)(DMF)₂ were determined by single-crystal XRD and supported by DFT simulations, respectively.²⁶ The structure of the activated Fe₂(DSBDC) is proposed based on its PXRD pattern. Green, yellow, red, blue, grey, and white spheres represent Fe, S, O, N, C, and H atoms, respectively. Unbound DMF molecules and framework H atoms are omitted for clarity.

**Fig. 2** IR spectra of various forms of Fe₂(DSBDC). Characteristic peaks of DMF, MeOH, and DCM are highlighted.
formula that matched that of fully desolvated Fe₂(DSBDC), whose IR spectrum also indicates the complete absence of DMF, MeOH, or THF characteristic bands (Fig. 2 and ESI†). Most tellingly, an N₂ adsorption of Fe₂(DSBDC) exhibits a type I isotherm with a saturation uptake of approximately 160 cm³ of N₂ per g at 77 K (Fig. 4) and an apparent Brunauer–Emmett–Teller (BET) surface area of 624 m² g⁻¹ (211 m² mmol⁻¹) (Fig. S4†). Although smaller than the value observed for related Mn₂(DSBDC) (329 m² mmol⁻¹)³⁵ and MOF-74 analogues (287–416 m² mmol⁻¹),³⁰,³²,³³ possibly due to a further structural distortion (vide infra), the apparent surface area of Fe₂(DSBDC) is significantly larger than that of Fe₂(DSBDC)(DMF)₂ (83 m² g⁻¹, 40 m² mmol⁻¹) (Fig. 4),³⁶ as would be expected upon removal of pore-blocking DMF molecules.

Although Fe₂(DSBDC) is crystalline, its PXRD pattern diverges from those of both Fe₂(DSBDC)(DMF)₂ and Fe₂(DSBDC)(DMF)₂: peaks at 6.6° and 11.3° corresponding to the [2–10] and [300] planes in the original regular hexagonal lattice are intact, but a new peak at 10.5° appears (Fig. 3). Qualitatively, this confirms that the hexagonal symmetry and pore shape is retained in Fe₂(DSBDC), but the lattice suffers a distortion along the crystallographic c direction, which runs parallel to the (–Fe–S–)₅ chains. Unfortunately, the relatively low crystallinity of the activated material does not offer sufficient quality for further structural refinement. However, soaking Fe₂(DSBDC) in DMF or DCM produces materials whose PXRD patterns are identical to that of original Fe₂(DSBDC)(DMF)₂:x(DMF) (Fig. 3),‡ This confirms that the distortion associated with solvent exchange and evacuation does not affect the connectivity of the framework. Similarly, any changes in electrical properties stemming from these distortions should also be reversible.

The electrical properties of Fe₂(DSBDC) were investigated with two-contact-probe devices in air-free conditions at 297 K using pressed pellets made by in situ press, an apparatus that was introduced previously.³⁷,³⁸ Single-crystal electrical conductivity measurements are unfeasible for all phases of Fe₂(DSBDC) because of the small crystallite size and extreme sensitivity to air. As shown in Fig. 5, current–voltage curves of all samples are linear, allowing the use of Ohm’s Law for resistance calculations. Notably, these measurements revealed that the electrical conductivity increases with the degree of solvation: it is 3.9 × 10⁻⁶ S cm⁻¹ for Fe₂(DSBDC)(DMF)₂:x(DMF), it drops to 5.8 × 10⁻⁷ S cm⁻¹ for Fe₂(DSBDC)(DMF)₂ upon removal of unbound DMF, and decreases further upon complete removal of DMF to only 1.5 × 10⁻⁹ S cm⁻¹ in fully desolvated Fe₂(DSBDC). In line with the reversible structural distortions discussed above, the electrical properties are also recovered upon soaking Fe₂(DSBDC) or Fe₂(DSBDC)(DMF)₂ in DMF.
with the recovery values of 3.1 × 10^{-7} \text{ S cm}^{-1} \text{ and } 2.3 \times 10^{-6} \text{ S cm}^{-1}, \text{ respectively}, \text{ in good agreement with the conductivity of pristine, as-synthesized Fe}_2(\text{DSBDC})(\text{DMF})_2 \times (\text{DMF}). \text{ (Scheme S1)}.

It should be noted, however, that coordination to the Fe center is critical for improved electrical conductivity. Indeed, although soaking fully desolvated Fe\textsubscript{2}(DSBDC) in DCM recovers the skeleton structure observed in Fe\textsubscript{2}(DSBDC) (DMF)\textsubscript{2} \times (DMF) (Fig. 3), DCM treatment does not recover electrical conductivity, which remains at 7.3 × 10^{-7} \text{ S cm}^{-1} \text{ (Fig. 5), similar to that of activated Fe\textsubscript{2}(DSBDC) itself. In other words, swelling with DCM (or presumably other non-coordinating solvents) recovers the original structure, but does not recover the electrical properties, which are more critically dependent on metal coordination rather than structural deformations.}

Density functional theory (DFT) calculations were further used to probe the origin of the coordination-induced electrical conductivity enhancement (Fig. 6). The crystal structure of the activated Fe\textsubscript{2}(DSBDC) was allowed to reach geometric and electronic equilibrium, resulting in regular hexagonal pores that resemble the structure of the DCM-soaked Fe\textsubscript{2}(DSBDC). The projected density of states (PDOS) shows that the valence band of activated Fe\textsubscript{2}(DSBDC) is dominated by Fe and S orbitals, suggesting that perturbation to either Fe or S valence will directly modify the valence band character. We hence conjecture that the major charge transport mechanism in activated Fe\textsubscript{2}(DSBDC) is hole hopping between Fe and S within the (-Fe-S-)\textsubscript{∞} chains, similar to the situation proposed in Fe\textsubscript{2}(DSBDC) (DMF)\textsubscript{2}.\textsuperscript{26,36}

Because the skeletons of Fe\textsubscript{2}(DSBDC)(DMF)\textsubscript{2} and the DCM-soaked Fe\textsubscript{2}(DSBDC) are isostructural, DCM likely does not affect charge mobility. Instead, it modulates charge density in the skeleton of the framework. Indeed, both a comparison of solid-state and truncated cluster calculations show that the work function of the activated Fe\textsubscript{2}(DSBDC) is smaller than the ionization potential of DMF by 1 eV (Fig. 6). As a result, partial electron transfer occurs when DMF binds to Fe centers in activated Fe\textsubscript{2}(DSBDC), as evidenced by approximately 1% reduction in local electron density of Fe, observed in both the solid-state and cluster calculations.\textsuperscript{§} Electron transfer thus generates holes as charge carriers, induces effectively mixed valency in (-Fe-S-)\textsubscript{∞} chains, and improves charge density and electrical conductivity.\textsuperscript{36}

**Conclusions**

DMF binding to the coordinatively unsaturated Fe centers in Fe\textsubscript{2}(DSBDC) leads to thousand-fold increase in the electrical conductivity at 297 K. DFT calculations identify electron transfer from Fe centers to the bound DMF molecules as a possible mechanism for improving charge density in the skeleton of the MOF. These results reinforce the importance of redox-matching between the framework and guest molecules to achieve electrically conductive MOFs, and point out the possibility of tuning electrical conductivity via coordinating guest molecules as well as applying MOFs with open metal sites for chemiresistive sensing of coordinating molecules.

**Conflicts of interest**

There are no conflicts to declare.

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**Notes and references**

\textsuperscript{†} The structural reversibility and crystallinity of activated Fe\textsubscript{2}(DSBDC) rule out desolvation-induced local destruction because it tends to be irreversible and leads to amorphization.

\textsuperscript{§} The calculation results of Fe\textsubscript{2}(DSBDC)(DMF)\textsubscript{2}, reported in ref. 26 were used to analyze the electron density on Fe centers in Fe\textsubscript{2}(DSBDC)(DMF)\textsubscript{2}.