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A tetrathiafulvalene salt of the nitrite (NO_2^-) anion: investigations of the spin-Peierls phase†

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Electrocrystallization of *ortho*-dimethyltetrathiafulvalene (*o*-DMTTF) in the presence of a nitrite (NO_2^-) anion affords a 2:1 phase, namely (*o*-DMTTF)₂NO₂, the first example of an isolated cation radical salt with NO_2^- . The *o*-DMTTF molecule organizes into strongly dimerized stacks that adopt a rare chessboard organization, associated with a strongly 1D structure. As a consequence of dimerization, the salt exhibits a semiconducting behavior ($\sigma_{\text{RT}} = 5 \text{ S cm}^{-1}$, $E_{\text{act}} = 0.1 \text{ eV}$). The bent NO_2^- anion is disordered on the inversion center but does not exhibit any anion-ordering transition. Combined SQUID magnetometry and electron paramagnetic resonance (EPR) experiments demonstrate the appearance of a spin-Peierls transition with $T_{\text{SP}} = 70 \text{ K}$. The temperature dependence of the spin gap and associated dimerization parameter δ across the spin-Peierls transition have been determined. The low-temperature narrow EPR line observed below T_{SP} is attributable to intrinsic magnetic defects localized on the *o*-DMTTF molecules themselves. The broadening of the EPR line at higher magnetic fields is associated with the disordered NO_2^- anions surrounding these magnetic defects, contributing to the inhomogeneity of their g factor.

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10th Anniversary Statement

J. Mater. Chem. C provides a unique forum to present our progress on molecular conducting materials, establishing a bridge between organic/coordination chemistry and solid-state physics, building on the competences of both fields to unravel the solid-state properties of radical species, with emphasis on not only electronic but also structural characteristics and their associated phase transitions. The legitimate search for converting a material into a device (optical, magnetic or electronic) does not hinder the journal to welcome more fundamental results, a very important point as the original chemical systems presented in *J. Mater. Chem. C* are often sources of inspiration to address other topics relevant to the sister journals *J. Mater. Chem. A* (energy and sustainability) and *J. Mater. Chem. B* (biology and medicine). We wish all the best to the journal on the occasion of this 10th anniversary and thanks all members of the editorial office for their efforts and commitment.

1. Introduction

The spin-Peierls transition is a second-order phase transition associated with crystal lattice distortion in quasi-one-dimensional

antiferromagnetic materials. Below the spin-Peierls temperature (T_{SP}), deformation appears and the antiferromagnetic exchange coupling within the magnetic chain is no longer uniform (J) but alternated (J_1 and J_2). A temperature-dependent energy gap $\Delta(T)$ then forms between the fundamental energy level and a quasi-continuum of energy levels. By lowering the temperature, the magnetic susceptibility tends towards zero. First predicted in 1970,^{1–3} this type of transition can take place upon cooling in uniform, antiferromagnetically coupled spin chains to stabilize the fundamental level of energy and lift its degeneracy. In 1975, the first observation of a spin-Peierls transition was made by Bray *et al.* on the $[\text{TTF}][\text{CuS}_4\text{C}_4(\text{CF}_3)_4]$ material below $T_{\text{SP}} = 12 \text{ K}$.^{4,5} Since then, spin-Peierls transitions have been mainly observed in organic conductors such as tetracyanoquinodimethane (TCNQ) salts⁶ or Fabre (TMTTF = tetramethyltetrathiafulvalene) salts,⁷ as well as certain inorganic compounds such as CuGeO_3 ⁸ or TiOX ($X = \text{Cl}$ or Br).^{9,10} In $(\text{TMTTF})_2\text{PF}_6$, it has been observed by the elastic

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scattering of neutrons¹¹ at $T_{\text{SP}} = 18$ K and in $(\text{TMTTF})_2\text{AsF}_6$ at $T_{\text{SP}} = 13$ K. Based on continuous-wave EPR measurements, we recently revealed that the spin dynamics in the low-temperature regime in Fabre salts was dominated by defects within spin-dimerized chains.¹² Such defects polarize the antiferromagnetically coupled spins in their vicinity, thereby leading to a finite, local alternating magnetization around the defect site, which can be described as a soliton, *i.e.*, a $\text{spin-}\frac{1}{2}$ quasi-particle comprising many correlated spins, pinned to the defect. Moreover, details of spin dynamics deep in this spin-Peierls phase were investigated by pulse ESR experiments, which revealed Rabi oscillations as signatures of coherent spin dynamics. However, there are limits in the possibilities of pulsed EPR measurements as no spin echo could be observed. Also, $(\text{TMTTF})_2\text{X}$ salts have low spin-Peierls transition temperatures (10–20 K), which limits the temperature study of relaxation and coherence times.

In this respect, a novel family of TTF derivatives, originally reported in 1983,¹³ is based on a non-centrosymmetric TTF, namely, *ortho*-dimethyltetrafulvalene, abbreviated as *o*-DMTTF here. The halide salts of *o*-DMTTF adopt a 2:1 stoichiometry,¹⁴ namely, $(\text{o-DMTTF})_2\text{X}$ (X = Cl, Br, I), with a striking quadratic structure (space group $I4_2d$; Fig. 1), characterized with uniform *o*-DMTTF stacks organized in a chessboard-like pattern, with a 90° rotation of the long molecular axis in the nearest-neighbor stacks. This original structure gives the salts a highly 1D character and a metallic behavior at room temperature (RT).

Their temperature–pressure phase diagrams have been established by Foury *et al.* up to 1.8 GPa,¹⁵ demonstrating the appearance of a spin-Peierls transition at a much higher temperature ($T_{\text{SP}} = 50$ K) than in $(\text{TMTTF})_2\text{PF}_6$ ($T_{\text{SP}} = 19$ K), which itself has the highest T_{SP} value of the $(\text{TMTTF})_2\text{X}$ series. In these *o*-DMTTF halides, the spin dynamics in the low-temperature regime is also dominated by solitons,¹⁶ and pulse ESR experiments in the spin-Peierls phase revealed Rabi oscillations as signatures of coherent spin dynamics. Further studies¹⁷ gave access to the temperature dependence of intra-chain coupling $J_{\text{eff}}(T)$, the temperature dependence of the dimerization parameter and the gap across the spin-Peierls

transition, and the content of spin-chain defects. We also showed that one-half of the total number of solitons are isolated (as singlets), whereas the other half forms pairs (soliton dimers) with a strong magnetic coupling.

Looking for other salts with a similar structure and potentially even higher SP transition temperatures, we investigated the nitrate (NO_3^-) salt, *i.e.*, $(\text{o-DMTTF})_2\text{NO}_3$.¹⁸ It crystallized in a lower-symmetry monoclinic space group ($P2_1/c$), preserving the chessboard-like pattern (Fig. 1) found in $(\text{o-DMTTF})\text{X}$ (X = Cl, Br, I). The stacks are, however, now strongly dimerized, and the NO_3^- anion is disordered on the inversion center, with the nitrogen atom at the center, and the anion rotating around the fixed N atom. This semiconducting salt exhibits two decoupled phase transitions, an anion ordering one at 124 K and a spin-Peierls one at 90 K, which is an original feature compared with TMTTF salts with non-centrosymmetric anions such as BF_4^- , ClO_4^- , and ReO_4^- , where both transitions are coupled and observed at the same temperature.

To potentially avoid the disorder problems and anion-ordering processes encountered with the NO_3^- anion, we considered the use of the smaller nitrite (NO_2^-) anion. Surprisingly, this anion has never been isolated in cation radical salts, perhaps owing to its known reducing character. For example, the electrogenerated perylene cation radical is known to instantaneously react with a nitrite ion in a 100% electron transfer process, yielding perylene and dinitrogen tetroxide.¹⁹ The reaction between the two latter species leads to the formation of mononitroperylene in excellent yields. Further, while working on similar naphthalene nitration, the electrochemical investigations of NO_2^- alone in CH_3CN showed that the anion oxidizes around +0.50 V *vs.* SCE,^{20,21} while perylene oxidizes at +0.85 V and naphthalene at +1.62 V. It follows that the cation radicals of organic donor molecules oxidizing below +0.5 V *vs.* SCE should be stable in the presence of NO_2^- . Indeed, we have found that acetonitrile solutions of tetrabutylammonium nitrite could be used as an electrolyte to successfully perform electrocrystallization experiments with TTF derivatives oxidizing below +0.5 V. Here, we report the structure of the 2:1 mixed-valence salt $(\text{o-DMTTF})_2\text{NO}_2$, together with its transport and magnetic properties. Inspired by the observation of solitons and soliton dimers in the spin-Peierls phase of quadratic halide salts with the same donor molecule $(\text{o-DMTTF})_2\text{X}$ (X = Cl, Br, I),^{16,17} by using EPR here, we comprehensively analyze the characteristics of the SP phase of $(\text{o-DMTTF})_2\text{NO}_2$, taking advantage of the very high transition temperature found in this salt at $T_{\text{SP}} = 70$ K.

2. Experimental section

2.1 Synthesis

Here, *o*-DMTTF was prepared as previously described²² and TBA nitrite was obtained from Aldrich and used as received. The electrocrystallization of *o*-DMTTF (10 mg) was performed in two-compartment cells with Pt electrodes (length: 15 mm; diameter: 1 mm) in dry CH_3CN (10 mL) containing TBA nitrite

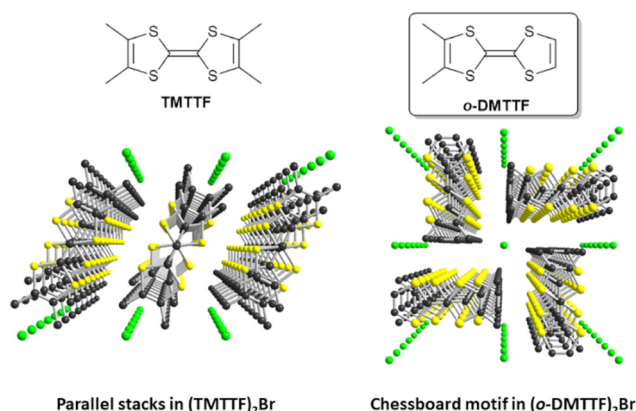


Fig. 1 Comparative solid-state organizations of the mixed-valence stacks in the Fabre salt $(\text{TMTTF})_2\text{Br}$ (left) and $(\text{o-DMTTF})_2\text{Br}$ (right).



(100 mg) as the electrolyte, at a constant current (1 μ A) and low temperature (5 $^{\circ}$ C). The crystals were harvested after ten days and washed with cold dry CH_3CN .

2.2 Structural characterization

A crystal of $(o\text{-DMTTF})_2\text{NO}_2$ suitable for X-ray diffraction single-crystal experiment was selected and mounted with a cryoloop on the goniometer head of a D8 Venture diffractometer equipped with a (CMOS) PHOTON 100 detector, using Mo- $\text{K}\alpha$ radiation ($\lambda = 0.71073$ $\text{\AA}$; multilayer monochromator). The structure was solved by a dual-space algorithm using the SHELXT program,²³ and then refined with full-matrix least-squares methods based on F^2 (SHELXL program).²⁴ All the non-hydrogen atoms were refined with anisotropic atomic displacement parameters. The H atoms were finally included in their calculated positions using the AFIX 137 procedure and treated as riding on their parent atom with constrained thermal parameters. Crystallographic data. $\text{C}_{16}\text{H}_{16}\text{NO}_2\text{S}_8$, $M = 510.78$, monoclinic, $a = 6.9491(6)$, $b = 12.3251(10)$, $c = 12.0663(9)$ $\text{\AA}$, $\beta = 103.318(3)^{\circ}$, $U = 1005.67(14)$ $\text{\AA}^3$, $T = 150(2)$ K, space group $P2_1/c$ (no. 14), $Z = 2$, 15795 reflections measured, 2295 unique ($R_{\text{int}} = 0.0434$), which were used in all the calculations. The final $wR(F^2)$ was 0.0386 (all data).

2.3 Magnetic susceptibility measurements

The magnetic measurements were performed with a SQUID magnetometer MPMS-XL from Quantum Design, in the temperature range from 2 to 300 K. A large number of single crystals were harvested from several electrochemical cells to reach 37.35 mg of $(o\text{-DMTTF})_2\text{NO}_2$ and enclosed in a Teflon pocket. The contribution from the Teflon bag as well as the diamagnetism of $(o\text{-DMTTF})_2\text{NO}_2$ have been subtracted. The temperature dependence of susceptibility has been performed under a magnetic field of 5 T.

2.4 Electron paramagnetic resonance

Electron paramagnetic resonance measurements have been performed using a conventional Bruker EMX spectrometer operating in the X band (about 9.39 GHz). A single crystal of $(o\text{-DMTTF})_2\text{NO}_2$ was glued on a Suprasil rod using Apiezon grease. The sample holder was attached to an automatic goniometer and the sample was centered in the cavity. The temperature was controlled using an Oxford ITC and ESR900 cryostat. The low-temperature measurements were performed using a cryogen-free Stinger system from ColdEdge, allowing to reach 7 K of base temperature.

2.5 Theoretical calculations

The tight-binding $\beta_{\text{HOMO-HOMO}}$ interaction energy calculations were based on the effective one-electron Hamiltonian of the extended Hückel method,²⁵ as implemented in the Caesar 1.0 chain of programs.²⁶ The off-diagonal matrix elements of the Hamiltonian were calculated according to the modified Wolfsberg-Helmholtz formula.²⁷ All the valence electrons were explicitly taken into account in the calculations and the basis set consisted of double- ζ Slater-type orbitals for all the atoms except for H (simple- ζ Slater-type orbital) using the Roothaan-Hartree-Fock wave functions of Clementi & Roetti.²⁸

All the other theoretical calculations were based on density functional theory (DFT) calculations and were performed with the ORCA program package.²⁹ Geometry optimization and electronic structure calculations were undertaken using the hybrid functional B3LYP^{30,31} in combination with the TZV/P basis set for all the atoms,³² and by taking advantage of the resolution of identity (RI) approximation in the Split-RI-J variant³³ with the appropriate Coulomb fitting sets.³⁴ Increased integration grids (Grid4 and GridX4 in ORCA convention) and tight SCF convergence criteria were used in the calculations.

3. Results and discussion

The electrocrystallization of $o\text{-DMTTF}$ was performed in CH_3CN at low temperatures (5 $^{\circ}$ C) as the salt proved to be too soluble at higher temperatures. The isolated crystals analyze as the 2 : 1 salt, i.e., $(o\text{-DMTTF})_2\text{NO}_2$. The preparation of this salt demonstrates that the nitrite anion can be effectively used as an electrolyte, at least with electron donors with a low oxidation potential, opening the way to a variety of other radical cation salts. Here, $(o\text{-DMTTF})_2\text{NO}_2$ crystallizes (at 150 K) in the monoclinic system, space group $P2_1/c$, with the $o\text{-DMTTF}$ molecule in the general position, a structure very close to that of NO_3^- salt.¹⁸ A projection view along the stacking axis (Fig. 2a) shows the general solid-state organization, with a 90° rotation of the long molecular axis in the nearest-neighbor stacks, affording this rare chessboard-like motif in TTF salts, already observed in the halide and nitrate salts of $o\text{-DMTTF}$.^{14,18} The NO_2^- anion adopts the expected bent structure due to the presence of an active lone pair on the nitrogen atom, with an O–N–O angle of $144.6(4)^{\circ}$ —a large value compared with those reported earlier.³⁵ This NO_2^- anion is disordered on the inversion center, with both oxygen atoms common to both orientations. A set of six C–H $\cdots$ O hydrogen bonds can be identified involving the CH_3 groups of $o\text{-DMTTF}$, with H $\cdots$ O distances of 2.669(3), 2.744(3), and 2.797(5) $\text{\AA}$ ($\times 2$), with the associated $\text{C}_{\text{Me}}\text{--H}\cdots\text{O}$ angles at $172.6(1)^{\circ}$, $165.6(1)^{\circ}$, and $151.3(1)^{\circ}$ (Fig. 2b). Despite a closely related structure, this NO_2^- salt differs from the reported NO_3^- salt where anion ordering was associated with the strengthening of such C–H $\cdots$ O hydrogen bonds. The disorder model adopted

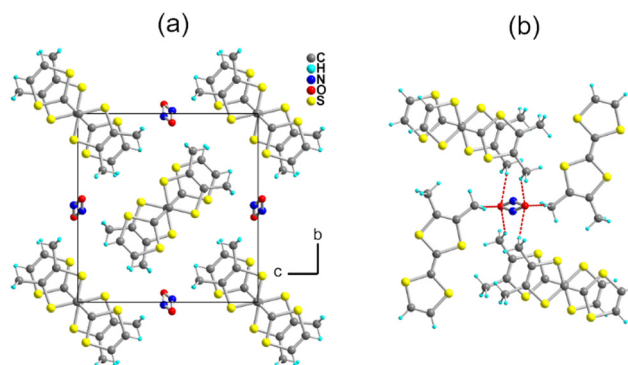


Fig. 2 (a) Projection view along a of the unit cell of $(o\text{-DMTTF})_2\text{NO}_2$. (b) Details of the C–H $\cdots$ O hydrogen bonds (red dotted lines).



here by the nitrite anion (at 150 K) lets us infer that a possible anion ordering would not change the atomic positions of the partially negatively charged oxygen atoms and hence would not significantly modify this C–H...O hydrogen bonding pattern at the organic/inorganic interface. An examination of the crystals at lower temperatures to tentatively observe the signature of an anion-ordering process was conducted on the CRYSTAL line at SOLEIL, above (85 K) and well below (35 K) the spin-Peierls transition ($T_{\text{SP}} = 70$ K). A comparison with the structure at 150 K showed no changes, with the same $P2_1/c$ space group, as well as the absence of any diffusion or substructure features.

The donor molecules stack along the a axis, with an alternation of intermolecular overlaps within the chain (Fig. 3a): a strong overlap of nearly eclipsed molecules and a weaker one with the so-called bond-over-ring overlap (Fig. 3b). This analysis is corroborated by the calculation of the $\beta_{\text{HOMO-HOMO}}$ overlap interaction energies, with $\beta_{\text{intra}} = 0.71$ and $\beta_{\text{inter}} = 0.39$ eV. The degree of alternation of the stack, defined as $2(\beta_{\text{intra}} - \beta_{\text{inter}})/(\beta_{\text{intra}} + \beta_{\text{inter}})$, is found at 0.58, a value comparable with that found in NO_3^- salt (0.60), but much stronger than those reported in TMTTF salts with PF_6^- (0.38) or Br^- (0.11).³⁶

Resistivity measurements (Fig. 4) performed on single crystals show an RT conductivity of 5 S.cm^{-1} and, upon cooling, an activated behavior with $\rho = \rho_0 \exp(E_{\text{act}}/kT)$ with $E_{\text{act}} = 0.1$ eV. This indicates that electron localization takes place with a gap of charge $\Delta_p \approx 2E_{\text{act}} \approx 0.2$ eV, a combined effect of the strong 1D character and electron–electron correlations associated with the strong stack alternation.

The temperature dependence of magnetic susceptibility $\chi(T)$ (Fig. 5a) has been evaluated using ESR and SQUID measurements. Here, $\chi(T)$ from ESR have been extracted from the spectra using eqn (1)

$$I_{\text{ESR}} = A \left(\frac{\Gamma \cos \varphi}{\Gamma^2 + (H - H_0)^2} + \frac{(H - H_0) \sin \varphi}{\Gamma^2 + (H - H_0)^2} \right) \quad (1)$$

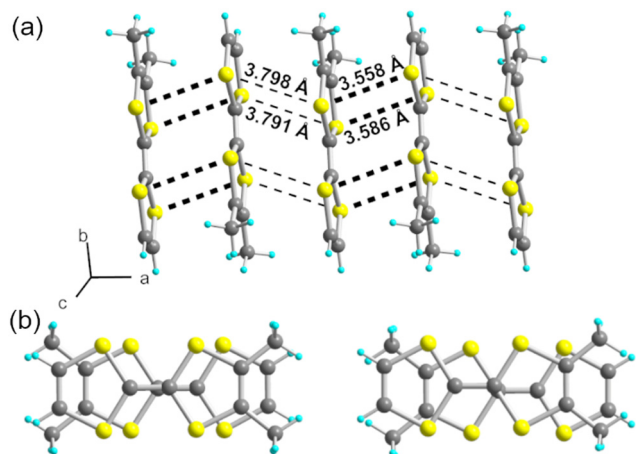


Fig. 3 (a) A side view of one chain in $(o\text{-DMTTF})_2\text{NO}_2$ (stacking axis a) with intermolecular S...S contacts shown as black dotted lines. (b) The two different overlap patterns within the stack: (left) intra-dyad and (right) inter-dyad (see main text).

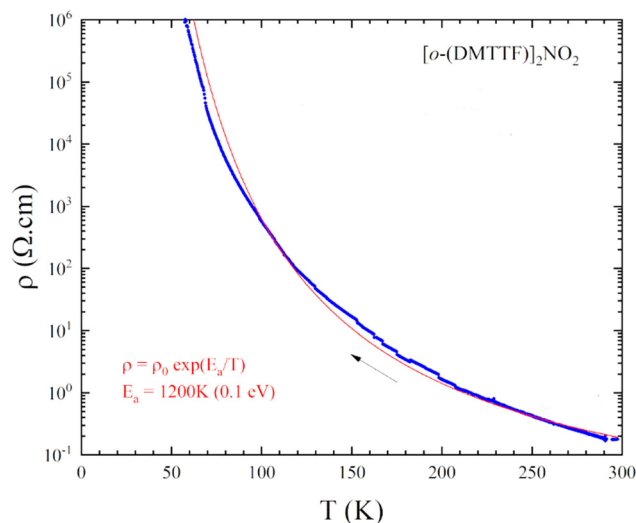


Fig. 4 Temperature dependence of the resistivity of $(o\text{-DMTTF})_2\text{NO}_2$.

where A is the amplitude of the ESR spectrum directly proportional to the magnetic susceptibility, Γ is the half-width at half-maximum, H_0 is the resonance field, and φ is the angle of dispersion. We use this fit procedure on our spectra for each temperature and each crystal orientation. A series of representative EPR spectra as a function of temperature are given in the ESI† (Fig S1)

In order to obtain quantitative ESR susceptibility, we renormalized it using SQUID measurement of the same compound at $T = 300$ K. In Fig. 5a, the squares denote SQUID data whereas circles denote the ESR data. A good agreement was found between the SQUID and ESR data down to 50 K. However, below 50 K, the susceptibilities differ significantly, and a paramagnetic signal following Curie behavior is visible. At lower temperatures, a concentration of impurities of 0.079% and 0.37% was extracted from the Curie behavior assuming $S = \frac{1}{2}$ and $g = 2$ from the ESR and SQUID data, respectively. Evidently, the quantity of impurities is clearly larger in SQUID measurements than in ESR. This difference can be attributed to the high selectivity of ESR measurements, which only provide information at a specific resonance frequency on a single crystal whereas SQUID magnetometry measures the total magnetic moment of a polycrystalline sample. Fig. 5b shows the spin-chain susceptibilities χ_c upon the removal of Curie contributions. Here, the SQUID and ESR data are identical for the full temperature range and within the limit of experimental error. Two behaviors are evident. From RT down to about 70 K, the susceptibility is weakly temperature dependent and can be described by a uniform-exchange Heisenberg spin chain. To quantitatively describe this part of susceptibility, it is common to use the well-known Bonner–Fisher curve.³⁷ This method assumes that the exchange coupling integral J is temperature independent, which is not the case here like in most organic spin chains. Indeed, the thermal variation in the crystallographic unit-cell parameters (Table S1, ESI†) shows that the cell contraction essentially affects the stacking axis a and thus



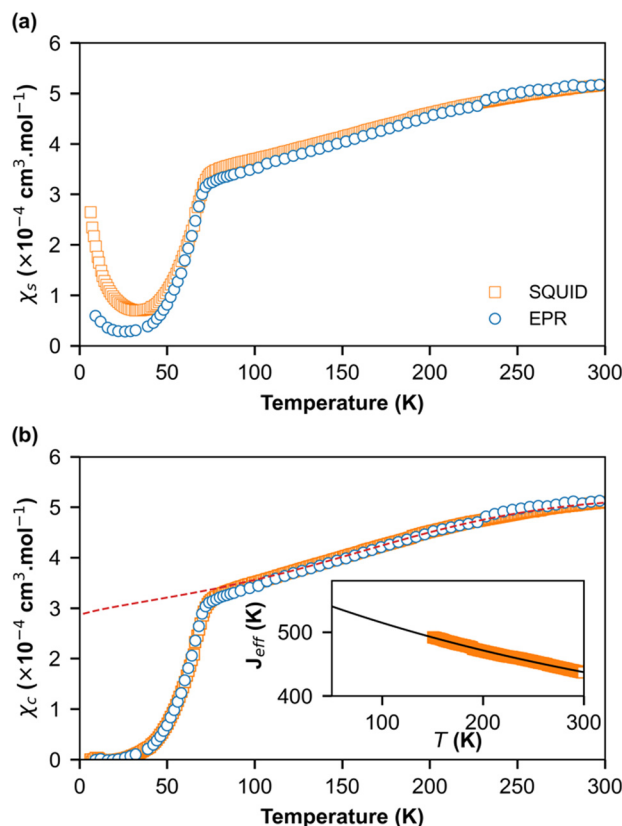


Fig. 5 (a) Temperature dependence of the spin susceptibility χ_s deduced from ESR (circles) and SQUID (square) in $(o\text{-DMTTF})_2\text{NO}_2$. (b) Temperature dependence of the spin susceptibility (χ_c) corrected for the Curie tail of $S = 1/2$ defects. The dashed lines are the theoretical values of susceptibility using exchange constants J_{eff} presented in the inset and extrapolated to lower temperatures.

the distance between the spins in chains, which decreases when T decreases. In order to extract the temperature dependence of J , we used a method developed by Johnston *et al.*³⁸ and successfully applied it to $(o\text{-DMTTF})_2\text{X}$ ($\text{X} = \text{Cl}, \text{Br}, \text{I}$).¹⁷ The temperature dependence of J is presented in the inset of Fig. 5b. We have limited the temperature range of extraction of J from 150 K to 300 K to avoid any effect of fluctuations near the phase transition. We then extrapolate to lower temperatures. Finally, we have simulated the susceptibility in the full temperature range using the extrapolated J (red dashed line in Fig. 5b).

Below $T_{\text{SP}} = 70$ K, the gapless Heisenberg uniform spin chain progressively enters into a non-magnetic gapped state through the spin-Peierls transition. The determination of spin gap Δ is imperative since it is related to the dimerization parameter δ . Since each spin is delocalized on an $o\text{-DMTTF}$ dyad, this dimerization parameter is associated with the tetramerization of $o\text{-DMTTF}$ stacks. However, the determination of an accurate value of Δ is tricky in spin-Peierls transitions, which are second-order transitions. Δ is temperature dependent close to T_{SP} and this dependence has to be considered. We have first estimated Δ in the temperature-independent regime (for $T < 0.5 T_{\text{SP}}$)³⁹

using the Bulaevski model⁴⁰ for the susceptibility of an alternated spin chain:

$$\chi_s(T) = N\alpha \frac{g^2 \mu_B^2}{k_B T} \alpha(\delta) \exp\left(-\frac{\Delta_B(\delta) J_1}{k_B T}\right)$$

where $J_1 = J(1 + \delta)$, α is the amplitude factor, and $\Delta_B J_1$ is the magnetic gap. We obtained $\delta_{(T=0)} = 0.115 \pm 0.005$ and $\Delta_{(T=0)} = 200 \pm 20$ K. For the temperature range where the gap is not stabilized, we had to consider the effect of temperature. Consequently, we used the Johnston *et al.* method³⁸ that individually fits each point. The temperature dependence of alternation parameter $\delta(T)$ and magnetic gap $\Delta(T)$ are given in Fig. 6.

Note that this method can be used only if the susceptibility is not too close to zero. Therefore, for $T < 30$ K, no value is provided. The results given by EPR and SQUID measurements are comparable. For $T < 40$ K, the gap and dimerization factor are temperature independent; consequently, as the temperature increases, δ and Δ decrease but do not vanish at $T = T_{\text{SP}}$ and above. This behavior is attributed to the existence of a pseudo-gap close but above T_{SP} , as already observed by X-ray diffraction studies in $(o\text{-DMTTF})_2\text{Br}$.¹⁵

Fig. 7 shows the temperature dependence of EPR linewidth $\Gamma(T)$ of $(o\text{-DMTTF})_2\text{NO}_2$. From 300 K down to $T_{\text{SP}} = 70$ K, the

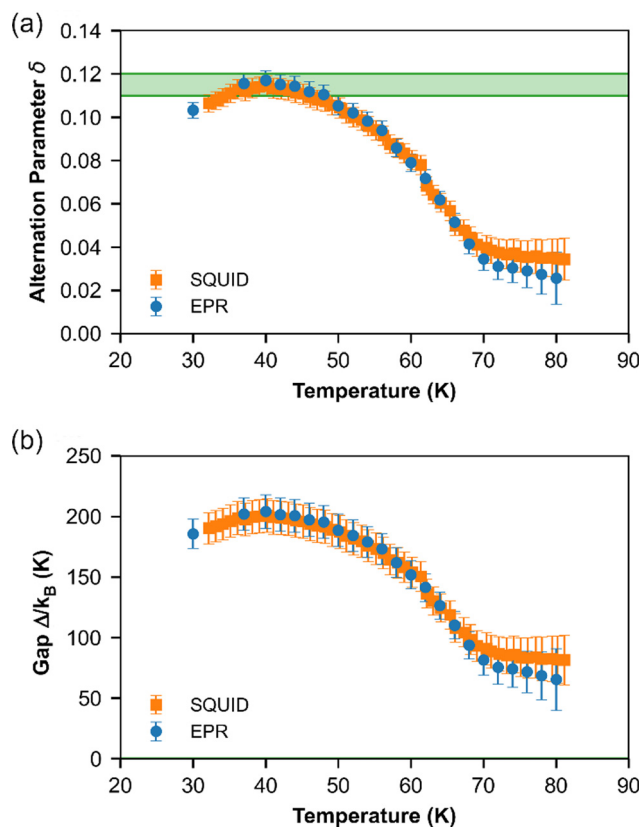


Fig. 6 Temperature dependence of (a) dimerization parameter δ and (b) spin gap Δ for $(o\text{-DMTTF})_2\text{NO}_2$ extracted from ESR (circle) and SQUID (square) measurements. The green zone represents the values estimated at $T = 0$ K with its error.

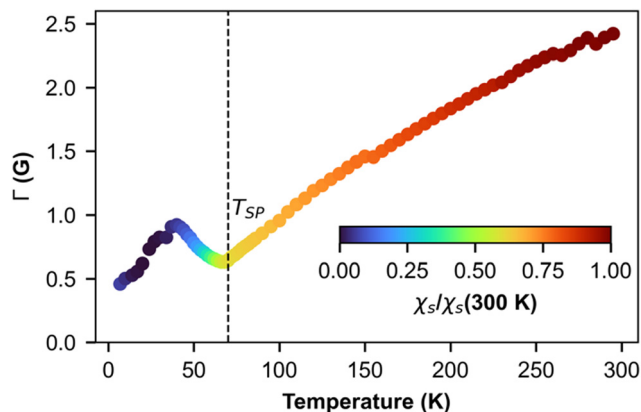


Fig. 7 Temperature dependence of ESR linewidth. The color map represents the intensity of the line presented in Fig. 4a.

linewidth decreases linearly as it is often observed in organic 1D conductors like Fabre salts $(\text{TMTTF})_2\text{X}$.⁴¹ On the contrary, in $(o\text{-DMTTF})_2\text{X}$ salts, the temperature dependence of the linewidth is completely different. $\Gamma(T)$ increases when T decreases for $\text{X} = \text{Cl}, \text{Br}, \text{I}$,¹⁵ while it decreases here for $\text{X} = \text{NO}_2$. Moreover, $\Gamma_{(T=T_{\text{SP}})}$ is much bigger for centrosymmetric halide anions (up to 100 G for $\text{X} = \text{I}$) than for the non-centrosymmetric one with $\Gamma_{(T=T_{\text{SP}})} = 0.5\text{G}$. In halide salts,¹⁵ the temperature dependence of Γ was possibly attributed to the existence of chains rotated by 90° in the crystal cell. However, this assumption cannot stand since with $\text{X} = \text{NO}_2$, the crystal structure is similar but the linewidth increases with T while it decreases for $\text{X} = \text{Cl}, \text{Br}$, and I . We think that the temperature dependence of linewidth in this high-temperature range is more related to the centro-/non-centrosymmetric nature of the counterion. However, in the absence of a microscopic model, we cannot go further in this analysis. Now, we analyze the temperature dependence of linewidth below T_{SP} . For T decreasing from T_{SP} , the linewidth increases to reach a local maximum at about $T = 40\text{ K}$ and then decreases. To explain these regimes, we calculated the temperature dependence of linewidth in the spin-Peierls phase using the method of moments for an odd and even number of spins in the chain (Fig. S3 in ESI†). The model is simple and suffers from size effects but qualitatively describes our measurements. For an even number of spins (which describes the full chain), the opening of the gap increases the linewidth but the intensity decreases exponentially (Fig. 2b), whereas for an odd number of spins (which corresponds to defects or end chains), the linewidth decreases but the intensity increases with a Curie-like behavior. We conclude that for $40\text{ K} < T < T_{\text{SP}}$, the linewidth is ruled by the full spin chain, whereas for $T < 40\text{ K}$, it describes the defects in the spin chain.

Finally, we focus on the low-temperature range ($T < 30\text{ K}$) where the susceptibility shows a Curie-like behavior (Fig. 5a). In this temperature range, the χ values obtained from the ESR and SQUID data are different. While SQUID measurements are not selective and contain the information from both extrinsic and intrinsic impurities, ESR—a spectroscopic technique—is very

selective. The angular variation in the g -factor at RT and 8 K (Fig. 8) are identical, indicating that the signal at lower temperatures is not associated with extrinsic defects but with the $o\text{-DMTTF}$ molecule in the stack.^{12,17,42}

We used this intrinsic defect's EPR line to probe the crystal structure around it by performing low- and high-frequency EPR measurements (Fig. 9). Indeed, in the presence of a homogeneous g factor, the linewidth should not vary with the external field, as experimentally observed, for example, in $(o\text{-DMTTF})_2\text{X}$ salts ($\text{X} = \text{Cl}, \text{Br}, \text{I}$).⁴³ On the other hand, in $(o\text{-DMTTF})_2\text{NO}_2$, the linewidth increases with frequency, indicating a pure Zeeman broadening effect. This behavior is usually observed in disordered or inhomogeneous systems. In the present case, $(o\text{-DMTTF})_2\text{NO}_2$ is a single crystal and no Zeeman broadening has been observed in analogous centrosymmetric salts $(o\text{-DMTTF})_2\text{X}$ ($\text{X} = \text{Br}, \text{Cl}, \text{I}$). Thus, the origin of this broadening can be explained by the non-centrosymmetric nature of NO_2^- counterion and its disorder in the crystal structure (Fig. 2). We have performed DFT calculations to estimate the g factors for many different orientations of NO_2^- (Fig. S4 in ESI†). Indeed, around the $o\text{-DMTTF}$ -centered magnetic defects, each NO_2^- has two equivalent orientations and each defect is surrounded by many NO_2^- ions, potentially contributing to the inhomogeneity of the g factor. This method has been used, for example, to probe the structure of dimethylammonium magnesium formate, where the DMA cation is disordered on three equivalent positions.^{44,45} The number of configurations quickly becomes large and time-consuming for DFT. We have, therefore, chosen two extreme configurations: all NO_2^- ions pointing to the $(o\text{-DMTTF})_2^+$ dyads and all NO_2^- ions pointing out of the $(o\text{-DMTTF})_2^+$ dyads. The details of the results are presented in the ESI† (Table S2); we found that the distribution of g factor, based on these extreme configurations, is contained within $\Delta g^{\text{DFT}} = 2.5 \times 10^{-3}$. Experimentally, we can extract the distribution of g factor from the linewidth and we found $\Delta g^{\text{exp}} = 1.2 \times 10^{-3}$, which is in good agreement with the DFT calculations.

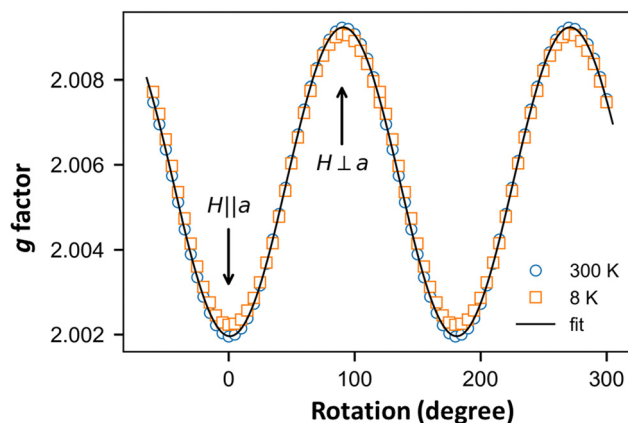


Fig. 8 Angular dependence of g factor of the EPR line attributed to defects in the spin chain measured at $T = 8\text{ K}$ (red squares) and of the uniform spin-chain line measured at $T = 300\text{ K}$ (green circles).



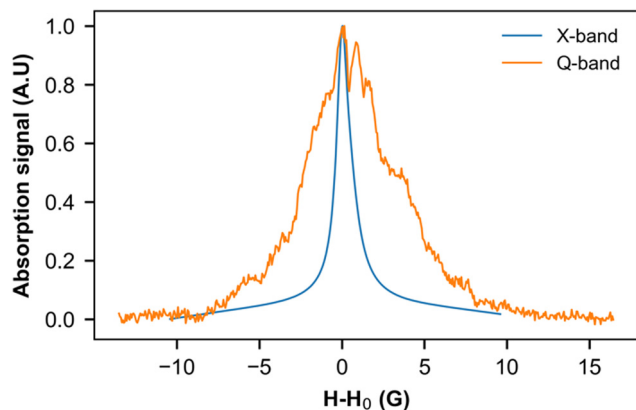


Fig. 9 EPR signal of intrinsic defects measured at 8 K in the X band (9.39 GHz) and Q band (33.75 GHz). To facilitate comparison, the resonance field H_0 has been subtracted from the magnetic field.

4. Conclusions

We have demonstrated here that radical cation salts of the nitrite anion can be stable in normal conditions and isolated as single crystals from the electrocrystallization of a TTF derivative with tetrabutylammonium nitrite as the electrolyte. Potentially, all the donor molecules whose oxidation potential falls below +0.5 V (vs. SCE) can now be considered for such experiments. In $(o\text{-DMTTF})_2\text{NO}_2$, the mixed-valence $o\text{-DMTTF}$ molecules organize into strongly dimerized stacks, with the stacks adopting a rare chessboard organization, associated with a strongly 1D structure. The bent, non-centrosymmetric NO_2^- anion is disordered on the inversion center, with the N atom on two symmetry-related positions (with 0.5 occupancy), whereas the two O atoms are not disordered. This peculiar disorder model and the prevalence of short $\text{C-H}\cdots\text{O}$ hydrogen bonds at the organic/inorganic interface explain the absence of an anion-ordering transition in this salt, contrary to those observed in other TTF salts with NO_3^- or ClO_4^- anions where the oxygens atoms, which are the ones acting as hydrogen bond acceptors, are disordered in the high-temperature phase whereas the N or Cl central atom is not.^{18,46} In accordance with the strongly dimerized stacks and 1D nature of the electronic structure, the $(o\text{-DMTTF})_2\text{NO}_2$ salt is a semiconductor with $\sigma_{\text{RT}} = 5 \text{ S.cm}^{-1}$ and $E_{\text{act}} = 0.1 \text{ eV}$. Combined SQUID and EPR experiments demonstrate the appearance of a spin-Peierls transition with $T_{\text{SP}} = 70 \text{ K}$. We have quantified the temperature dependence of the spin gap and associated dimerization parameter δ across the spin-Peierls transition and shown the existence of a pseudogap above T_{SP} . We also demonstrate that the low-temperature signal observed below T_{SP} is attributable to intrinsic magnetic defects, as observed earlier in $o\text{-DMTTF}$ halide salts, and that the broadening of the EPR line with the magnetic field can be associated with the disordered NO_2^- anions surrounding a magnetic defect, contributing to the inhomogeneity of the g factor.

The next step in the investigations of the magnetic properties of this salt will involve pulsed EPR experiments to explore

the dynamics of these low-temperature magnetic defects in the spin-Peierls phase, as reported in Fabre salts $(\text{TMTTF})_2\text{X}$ ($\text{X} = \text{PF}_6^-, \text{AsF}_6^-$)¹² and more recently in $o\text{-DMTTF}$ halide salts.¹⁷ In the latter, it was indeed shown that in the vicinity of such strongly correlated defects, the spins are antiferromagnetically polarized, forming a magnetic soliton: a spin- $\frac{1}{2}$ quasi-particle comprising many correlated spins, pinned to the defect. Such studies are simplified here because of the high spin-Peierls transition temperature (70 K) of $(o\text{-DMTTF})_2\text{NO}_2$ compared with Fabre salts (10–20 K). The outcome of these pulsed EPR experiments will be reported in future studies.

Data availability

The complete data that support the findings of this study are openly available in Zenodo at <http://doi.org/10.5281/zenodo.7433815>.

Conflicts of interest

There are no conflicts to declare.

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References

- 1 D. B. Chesnut, *J. Chem. Phys.*, 1966, **45**, 4677–4681.
- 2 P. Pincus, *Solid State Commun.*, 1971, **9**, 1971–1973.
- 3 E. Pytte, *Phys. Rev. B: Solid State*, 1974, **10**, 4637–4642.
- 4 J. W. Bray, H. R. Hart, L. V. Interrante, I. S. Jacobs, J. S. Kasper, G. D. Watkins, S. H. Wee and J. C. Bonner, *Phys. Rev. Lett.*, 1975, **35**, 744–747.
- 5 I. S. Jacobs, J. W. Bray, H. R. Hart, L. V. Interrante, J. S. Kasper, G. D. Watkins, D. E. Prober and J. C. Bonner, *Phys. Rev. B: Solid State*, 1976, **14**, 3036–3051.
- 6 S. Huizinga, J. Kommandeur, G. A. Sawatzky, B. T. Thole, K. Kopinga, W. J. M. de Jonge and J. Roos, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 1979, **19**, 4723–4732.
- 7 A. Maarou, S. Flandrois, G. Fillion and J. P. Morand, *Mol. Cryst. Liq. Cryst.*, 1985, **119**, 311–315.
- 8 M. Hase, I. Terasaki and K. Uchinokura, *Phys. Rev. Lett.*, 1983, **70**, 3651–3654.
- 9 V. Kataev, J. Baier, A. Möller, L. Jongen, G. Meyer and A. Freimuth, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2003, **68**, 140405.
- 10 D. V. Zakharov, J. Deisenhofer, H.-A. Krug von Nidda, P. Lunkenheimer, J. Hemberger, M. Hoinkis, M. Klemm,



- M. Sing, R. Claessen, M. V. Eremin, S. Horn and A. Loidl, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2006, **73**, 094452.
- 11 P. Foury-Leylekian, D. Le Bolloc'h, B. Hennion, S. Ravy, A. Moradpour and J.-P. Pouget, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2004, **70**, 180405.
 - 12 S. Bertaina, C.-E. Dutoit, J. Van Tol, M. Dressel, B. Barbara and A. Stepanov, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2014, **90**, 060404.
 - 13 A. Abderraba, R. Laversanne, E. Dupart, C. Coulon, P. Delhaes and C. Hauw, *J. Phys. Colloques*, 1983, **44**, C3, 1243–1246.
 - 14 M. Fourmigué, E. W. Reinheimer, K. R. Dunbar, P. Auban-Senzier, C. Pasquier and C. Coulon, *Dalton Trans.*, 2008, 4652–4658.
 - 15 P. Foury-Leylekian, P. Auban-Senzier, C. Coulon, O. Jeannin, M. Fourmigué, C. Pasquier and J.-P. Pouget, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2011, **84**, 195134.
 - 16 J. Zeisner, O. Pilone, L. Soriano, G. Gerbaud, H. Vezin, O. Jeannin, M. Fourmigué, B. Büchner, V. Kataev and S. Bertaina, *Phys. Rev. B*, 2019, **100**, 224414.
 - 17 L. Soriano, O. Pilone, M. D. Kuz'min, H. Vezin, O. Jeannin, M. Fourmigué, M. Orio and S. Bertaina, *Phys. Rev. B*, 2022, **105**, 064434.
 - 18 O. Jeannin, E. W. Reinheimer, P. Foury-Leylekian, J.-P. Pouget, P. Auban-Senzier, E. Trzop, E. Collet and M. Fourmigué, *IUCrJ*, 2018, **5**, 361–372.
 - 19 L. Eberson and F. Radner, *Acta Chem. Scand., Ser. B*, 1985, **39**, 357–374.
 - 20 M. N. Cortona, N. Veltorazzi, J. J. Silber and L. Sereno, *J. Electroanal. Chem.*, 1995, **394**, 245–251.
 - 21 T. L. Broder, D. S. Silvester, L. Aldous, C. Hardacre and R. G. Compton, *J. Phys. Chem. B*, 2007, **111**, 7778–7785.
 - 22 F. Gerson, A. Lamprecht and M. Fourmigué, *J. Chem. Soc., Perkin Trans. 2*, 1996, 1409–1414.
 - 23 G. M. Sheldrick, *Acta Crystallogr.*, 2015, **A71**, 3–8.
 - 24 G. M. Sheldrick, *Acta Crystallogr.*, 2015, **C71**, 3–8.
 - 25 M.-H. Whangbo and R. Hoffmann, *J. Am. Chem. Soc.*, 1978, **100**, 6093–6098.
 - 26 J. Ren, W. Liang and M.-H. Whangbo, *Crystal and Electronic Structure Analysis Using CAESAR*, PrimeColor Software Inc., 1998.
 - 27 J. Ammeter, H.-B. Bürgi, J. Thibeault and R. Hoffmann, *J. Am. Chem. Soc.*, 1978, **100**, 3686–3692.
 - 28 E. Clementi and C. Roetti, *At. Data Nucl. Data Tables*, 1974, **14**, 177–478.
 - 29 F. Neese, *Wiley Interdiscip. Rev.: Comput. Mol. Sci.*, 2012, **2**, 73–78.
 - 30 A. D. Becke, *J. Chem. Phys.*, 1993, **98**, 1372–1377.
 - 31 C. Lee, W. Yang and R. G. Parr, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 1988, **37**, 785–789.
 - 32 A. Schäfer, C. Huber and R. Ahlrichs, *J. Chem. Phys.*, 1994, **100**, 5829–5835.
 - 33 F. Weigend, *Phys. Chem. Chem. Phys.*, 2006, **8**, 1057–1065.
 - 34 A. Klamt and G. Schüürmann, *J. Chem. Soc., Perkin Trans. 2*, 1993, 799–805.
 - 35 A search in the CSD on other simple organic salts of the NO₂[−] anion shows a wide range of O–N–O angles between 103 and 141°, as found in BEWGUH (109.26°), IZUGUF (103.73°), KUCPIH (117.20°), KUHNK (116.84°), MAKRAR (115.10°), TEZVOJ (114.77°), VAJRIH (123.91°), YAWHIP (116.47°) and ZEMBUO (141.09°).
 - 36 J.-P. Pouget and S. Ravy, *J. Phys. I*, 1996, **6**, 1501–1525.
 - 37 J. C. Bonner and M. E. Fisher, *Phys. Rev.*, 1964, **135**, 640–658.
 - 38 D. C. Johnston, R. K. Kremer, M. Troyer, X. Wang, A. Klümper, S. L. Bud'ko, A. F. Panchula and P. C. Canfield, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2000, **61**, 9558–9606.
 - 39 E. Orignac and R. Chitra, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2004, **70**, 1–12.
 - 40 L. Bulaevskii, *Sov. Phys.-Solid State*, 1969, **11**, 1132.
 - 41 C. Coulon and R. Clérac, *Chem. Rev.*, 2004, **104**, 5655–5687.
 - 42 L. Soriano, J. Zeisner, V. Kataev, O. Pilone, M. Fourmigué, O. Jeannin, H. Vezin, M. Orio and S. Bertaina, *Appl. Magn. Reson.*, 2020, **51**, 1307–1320.
 - 43 L. Soriano, unpublished results.
 - 44 M. Orio, J. K. Bindra, J. van Tol, M. Giorgi, N. S. Dalal and S. Bertaina, *J. Chem. Phys.*, 2021, **154**, 154201.
 - 45 S. Bertaina, N. Abhyankar, M. Orio and N. S. Dalal, *J. Phys. Chem. C*, 2018, **122**, 16431–16436.
 - 46 J.-P. Pouget, *Crystals*, 2012, **2**, 466–520.

