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Synthesis and size-dependent spin crossover of coordination polymer [Fe(Htrz)₂(trz)](BF₄)[†]

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The synthesis of quality single crystals is central to materials chemistry for optical, magnetic, and electronic device applications. The present work reports on the synthesis of single crystals of iron-triazole coordination polymer [Fe(Htrz)₂(trz)](BF₄) where (Htrz) = 1*H*-1,2,4-triazole. Crystals of size as long as 80 μm can be achieved by controlling the temperature, precursor concentration, and solvent type. It is found that its thermal spin crossover depends largely on the crystal size. Fine crystals are ideal for depositing a thin film that exhibits redox activity. The largest crystals allow reliable electrical conductance measurements that reveal two different activation energies at the low spin state and the high spin state, which are one order of magnitude smaller than the electronic gaps calculated based on density functional theory. The synthetic route sought in the present study can be applied to other coordination polymers and related materials and provides the basis for their applications.

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1 Introduction

Spin crossover or spin transition is a phenomenon wherein a molecular complex changes its form from a low-spin (LS) state to a high-spin (HS) state under the influence of external factors such as temperature, pressure or external stimuli.¹ This very attractive phenomenon occurs in octahedral complexes of 3d¹–3d⁷ first row transition metal ions; Cr(II), Mn(III), Fe(III), Mn(II), Fe(II), Co(III) and Co(II). They are stable in both LS and HS states and thus, the two states co-exist at equilibrium. This gives rise to molecular bistability accompanied by thermal hysteresis and chromism, which provides the impetus for technological applications such as data storage and memory devices² and display devices.³ The spin crossover can also be brought about by light irradiation *i.e.* light-induced excited spin state trapping (LIESST) and reverse LIESST. This was successfully implemented in optical data storage and processing devices.^{4–6} Spin-crossover materials are central to some of the neat technologies such as pressure-controlled conductivity (piezoelectric effect)⁷ photoconduction, molecular switches, memory devices, *etc.*^{8–10} The development in the field of supramolecular structures,

covalent organic frameworks, molecular organic frameworks, and their tendency to exhibit spin crossover^{11–15} have revived the research in this field.

The present work aims to understand the synthesis and size-dependent spin crossover of a Fe(II) coordination polymer, [Fe(Htrz)₂(trz)](BF₄) (see Fig. 1). In [Fe(Htrz)₂(trz)](BF₄), Fe(II) having six electrons in the 3d shell is coordinated to three pairs of nitrogen atoms of 1,2,4-triazole in an octahedral geometry. The 3d orbitals are split to the t_{2g} and e_g levels (see Fig. 2). In the LS state, the ligand field stabilizing energy is greater than the pairing energy, hence the electrons pair-up to fully occupy the t_{2g}

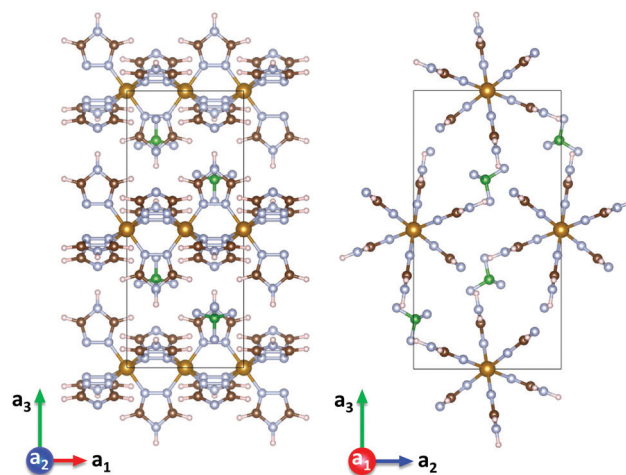


Fig. 1 Crystal structure of [Fe(Htrz)₂(trz)](BF₄) viewed along the a₂ axis (left) and along the a₁ axis (right), reproduced from structure files.¹⁶

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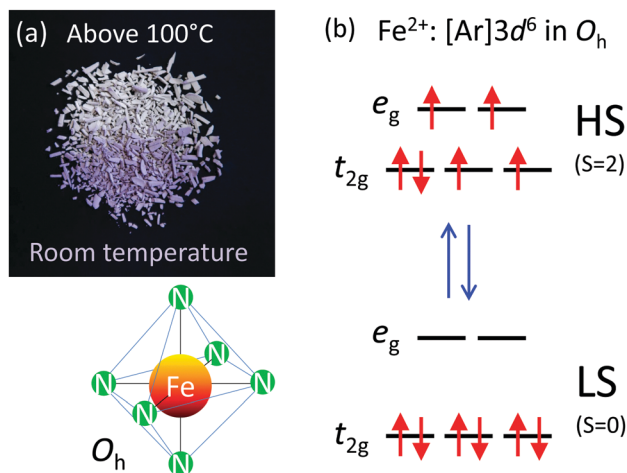


Fig. 2 (a) Spin crossover of $[\text{Fe}(\text{Htrz})_2(\text{trz})](\text{BF}_4)$ accompanied by the colour change from purple in the LS state at room temperature to white in the HS state at a temperature above 100 °C. (b) spin transition in $\text{Fe}^{2+}: [\text{Ar}]3d^6$ in an octahedral ligand geometry.

level, whereas in the HS state the ligand splitting is smaller as compared to the pairing energy, hence the electrons occupy the d orbitals according to Hund's rule of maximum multiplicity. Upon a LS-to-HS transition, two electrons are transferred from the t_{2g} subshell to the e_g subshell as the Fe-N bond length increases.^{8,10,13}

A series of systematic synthesis is presented to realise the dependence of the crystal size on the precursor concentration, temperature, and the type of solvent. It is found that much larger crystals are synthesised in water than in methanol. The length of crystalline needles can become as long as 80 μm by introducing a temperature gradient.

The phenomenon of spin crossover is accompanied by a change in color on heating, referred to as thermochromism. The color of $[\text{Fe}(\text{Htrz})_2(\text{trz})](\text{BF}_4)$ changes from purple in the LS state to white in the HS state (see Fig. 2a and Video S1, ESI†), which is attributed to spectral changes in UV-vis absorption.

Raman spectroscopy shows vibrational modes characteristic for the reported structure and their changes upon spin crossover as the temperature is increased from room temperature to 375 K. It is demonstrated that both spin transition temperature and hysteresis increase monotonically as the crystal size increases.

Drop-casting nanocrystalline powder in methanol can form homogeneous thin films with excellent adhesion to FTO-coated glass. The electrochemical analysis elucidates their redox activity which signifies its potential in a range of electronic and energy applications.

Finally, recent studies reported decent electrical conductance of nano- and micro-crystalline $[\text{Fe}(\text{Htrz})_2(\text{trz})](\text{BF}_4)$ that exhibited thermal hysteresis coinciding with their magnetic hysteresis. Our homogeneously long crystals are large enough to bridge standard source-drain electrodes that allow reliable measurements. We demonstrate that the temperature dependence of the electrical conductance follows the Arrhenius equation with an activation energy of 0.12 eV (0.24 eV) below (above) 384 K with no thermal

hysteresis. The activation energies are one order of magnitude smaller than the energy gaps calculated based on the density functional theory taking the derivative discontinuity of the exchange-correlation energy into account, indicating a conduction mechanism defined by the carrier injection barrier or defect levels.

2 Results and discussion

2.1 Synthesis

A systematic series of synthesis in water has been carried out in order to realise crystals of different dimensions and dependencies on the precursor concentration (C) and the temperature (T). A molar ratio of $[\text{Fe}(\text{BF}_4)_2 \cdot 6\text{H}_2\text{O}]:[\text{triazole}] = 1:3$ was maintained in the entire synthesis. Fig. 3 shows the optical micrographs of crystals prepared at $T = 45$ °C with different precursor concentrations in the range $C = 0.2$ – 0.4 M of $\text{Fe}(\text{BF}_4)_2 \cdot 6\text{H}_2\text{O}$. No large crystals are grown at concentrations out of the range presented. It is observed that the crystalline needles become thicker as the concentration is increased. At 45 °C, the length extends as the concentration is increased from $C = 0.2$ to 0.25 M, and then it gets shorter as the concentration is increased further to 0.4 M. With $C = 0.5$ M or higher no large crystals are grown. The size homogeneity tends to degrade by increasing the concentration.

Fig. 4 shows the crystals synthesised with the concentration fixed at $C = 0.3$ M in the temperature range of 50–70 °C. The crystalline needles became longer and thicker with the increasing temperature up to 60 °C. At higher temperatures ($T = 70$ °C) the crystals become much smaller. Furthermore, the crystals grow as large as 80 μm (see Fig. 5) with a temperature gradient

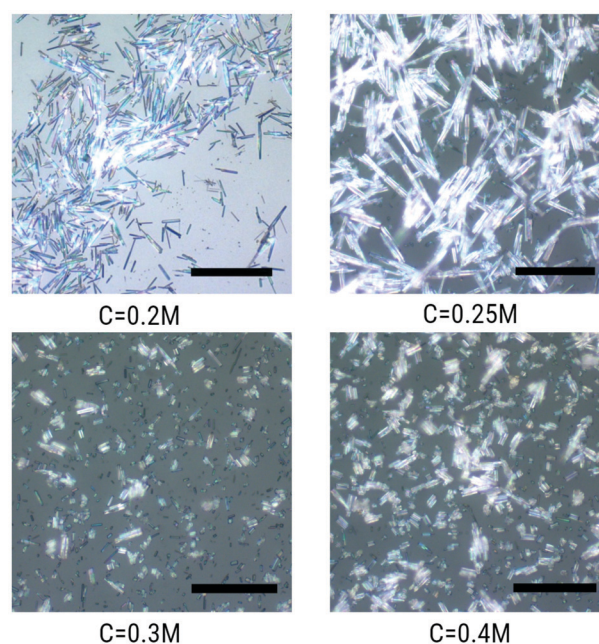


Fig. 3 Optical micrographs of $[\text{Fe}(\text{Htrz})_2(\text{trz})](\text{BF}_4)$ crystals synthesized in water at 45 °C with $\text{Fe}(\text{BF}_4)_2 \cdot 6\text{H}_2\text{O}$ concentrations of 0.2, 0.25, 0.3 and 0.4 M in water. Scale bars correspond to 50 μm .



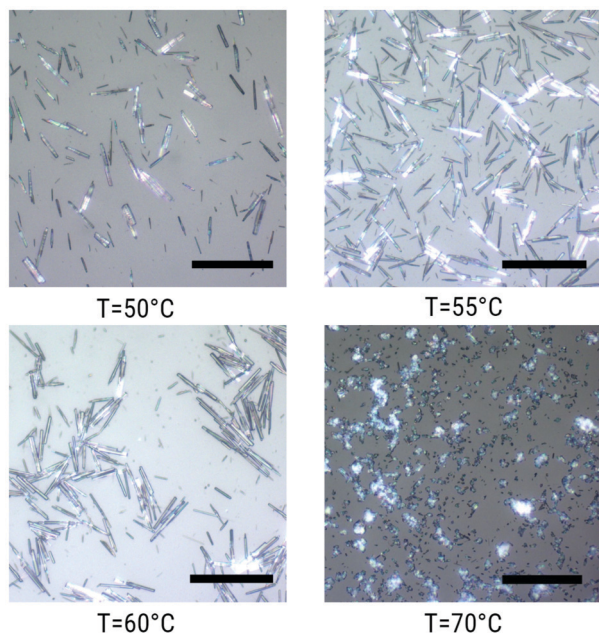


Fig. 4 Optical micrographs of $[\text{Fe}(\text{Htrz})_2(\text{trz})](\text{BF}_4)$ crystals synthesized in water at 50, 55, 60 and 70 °C with a $\text{Fe}(\text{BF}_4)_2 \cdot 6\text{H}_2\text{O}$ concentration of 0.3 M. Scale bars correspond to 50 μm .

around 45 °C (for more details, see the Methods section). To the best of our knowledge, no such large crystals of $[\text{Fe}(\text{Htrz})_2(\text{trz})](\text{BF}_4)$ were previously synthesized. To summarise, the average crystal size depends largely on the reaction temperature and precursor concentration. It is found that the largest crystals are grown homogeneously at $C = 0.3$ M and 60 °C. By introducing a temperature gradient, the crystals can grow as long as 80 μm .

2.2 X-ray diffraction

The powder X-ray diffraction profiles of the well-crystalline sample (crystal) prepared at $C = 0.3$ M and 45 °C and the

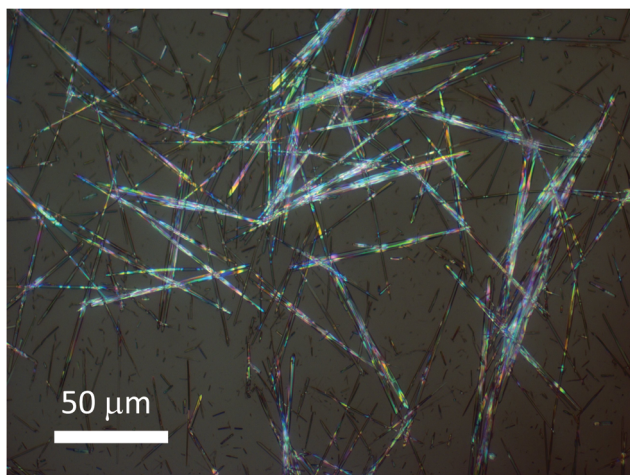


Fig. 5 Optical micrograph of $[\text{Fe}(\text{Htrz})_2(\text{trz})](\text{BF}_4)$ crystals synthesized in water with a temperature gradient around 45 °C with a $\text{Fe}(\text{BF}_4)_2 \cdot 6\text{H}_2\text{O}$ concentration of 0.3 M.

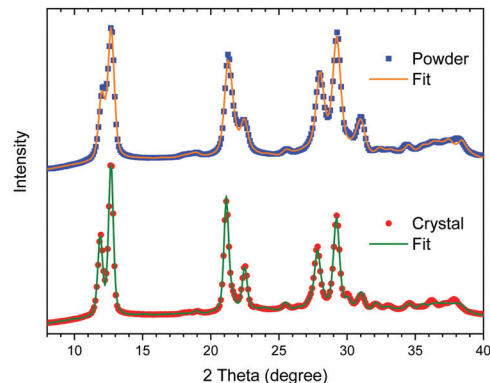


Fig. 6 Powder X-ray diffraction profiles of the well-crystalline (crystal) and least-crystalline (powder) $[\text{Fe}(\text{Htrz})_2(\text{trz})](\text{BF}_4)$ and the best-fits after Rietveld refinement.

least-crystalline sample (powder) synthesized in methanol are plotted in Fig. 6. Both diffraction profiles exhibit all peaks characteristic of the LS state of $[\text{Fe}(\text{Htrz})_2(\text{trz})](\text{BF}_4)$ reported previously,^{16–22} but not of $[\text{Fe}(\text{Htrz})_3](\text{BF}_4)$ which was often reported as an impurity.^{18,20} The lattice parameters obtained after Rietveld refinement match well with previously reported data.^{16,18} See ESI† for further details. The diffraction peaks of the powder are broader than those of the crystal, which can be attributed to much smaller crystallites. For detailed analysis, see ESI.†

2.3 UV-vis spectroscopy

The UV-vis absorption spectra of the powder sample in the LS and HS states are compared in Fig. 7. The absorption peak centred at 2.3 eV^{2,23} disappears upon the transition from the LS state to the HS state. This changes the sample colour from purple to white (see Fig. 2a) and leads to a larger optical gap.

2.4 Raman spectroscopy

Changes in the structure upon spin crossover can be probed by Raman spectroscopy. Fig. 8 shows the Raman spectra of the

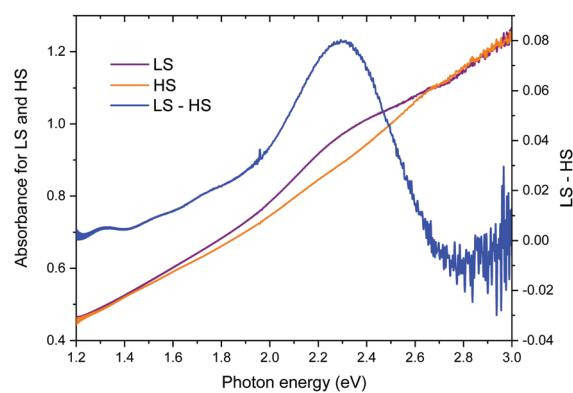


Fig. 7 UV-vis absorption spectra of the powder $[\text{Fe}(\text{Htrz})_2(\text{trz})](\text{BF}_4)$ in the LS and HS states. The difference spectrum (LS–HS) elucidates the absorption band centred at 2.3 eV in the LS state.



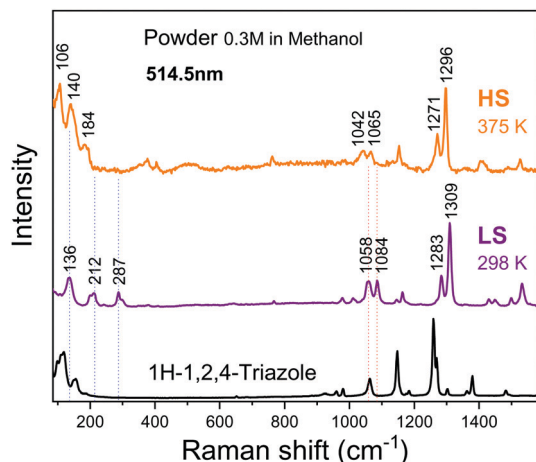


Fig. 8 Raman spectra of the powder sample at 298 K (LS state) and 375 K (HS state), measured at a laser wavelength of $\lambda = 514.5$ nm.

powder sample prepared in methanol measured at 298 and 375 K. Upon heating, the peaks at 136, 212 and 287 cm^{-1} indicated by the blue dashed vertical lines redshift to 106, 140 and 184 cm^{-1} , respectively, similar to those previously reported in the literature.¹⁸ The band at 287 cm^{-1} corresponds to a well-defined Fe–N stretching vibrational mode.²⁴ In the high-frequency region beyond 800 cm^{-1} , the peak shifts are minor, but noticeable as represented by the red short dot lines. $[\text{Fe}(\text{Htrz})_2(\text{trz})](\text{BF}_4)$ possess Htrz = 1H-1,2,4-triazole and trz = the deprotonated triazolato(–) ligands. The bands at 1058 and 1309 cm^{-1} of the LS state (1042 and 1296 cm^{-1} of the HS state) can be attributed to the Htrz and the bands at 1084 and 1283 cm^{-1} of the LS state (1065 and 1271 cm^{-1} of the HS state) to the trz.¹⁸ This excludes the presence of $[\text{Fe}(\text{Htrz})_3](\text{BF}_4)$ that was often reported as an impurity.^{18,20} All these changes are consistent with previous results^{18,24–32} and can be associated with the spin crossover. Upon a transition from the LS state to the HS state, the metal–ligand bond distance is extended as the ligand field is lowered and two electrons are transferred from the t_{2g} orbitals to the e_g orbitals. This leads to a lowering of metal–ligand stretching mode frequencies.^{18,24,26}

2.5 Magnetisation

In the present work, we compare the magnetic properties of the best-crystalline sample synthesised in water at $C = 0.3$ M and $T = 60$ °C and the powder sample synthesised in methanol at $C = 0.3$ M and room temperature. The corresponding magnetisation data in the temperature range of 320–400 K measured at 1 T are plotted in Fig. 9. The LS-to-HS transition occurs around $T_{1/2\downarrow} = 384$ K (360 K) upon heating while the reverse (HS-to-LS) transition occurs around $T_{1/2\uparrow} = 351$ K (338 K) upon cooling for the best-crystalline sample (the powder sample), where $T_{1/2}$ is the temperature at which 50% of molecules are in the LS state and 50% of molecules are in the HS state. Note that $T_{1/2\uparrow}$ is higher upon the first heating due to remanent solvent or moisture,^{33,34} as shown in the right panel of Fig. 9, so the third sweeps are compared in the left panel.

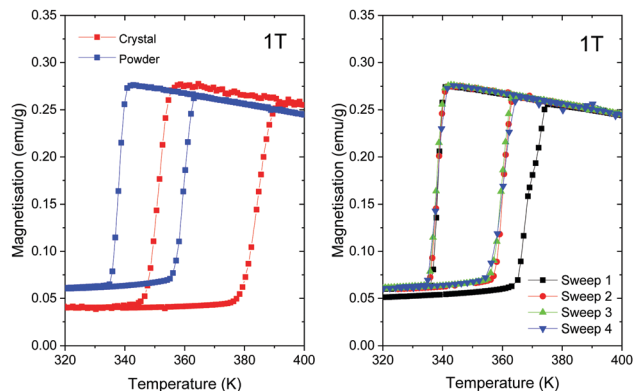


Fig. 9 (left) Temperature dependence of magnetisation (third sweep) for the well-crystalline (crystal) and least-crystalline (powder) $[\text{Fe}(\text{Htrz})_2(\text{trz})](\text{BF}_4)$ samples measured in a magnetic field of 1 T. (right) Various temperature sweep cycles for the powder sample, for details please see the text.

Both transition temperatures are higher for the best-crystalline sample. The corresponding width of the hysteresis loop is $\Delta T_{1/2} = 33$ K for the best-crystalline sample that is reduced to $\Delta T_{1/2} = 22$ K for the powder sample. The remanent magnetisation in the LS state, which can be attributed to high spin Fe(II) impurities, is about 15% of the maximum magnetization in the HS state for the best-crystalline sample and 22% for the powder sample. The fraction increases as the size of the nanoparticle decreases.

$T_{1/2\downarrow}$ for $[\text{Fe}(\text{Htrz})_2(\text{trz})](\text{BF}_4)$ synthesized under different conditions reported previously is in the range of 335–365 K, whereas $T_{1/2\uparrow}$ is in the range of 360–395 K.^{19,22,33–43} According to the previous studies on nanoparticles of different sizes,^{34,35,40,41} $T_{1/2\uparrow}$ is reduced as the volume of particles becomes smaller whereas $T_{1/2\downarrow}$ is less affected. Consequently, the hysteresis width is reduced for small particles. The transition temperatures of our powdery sample are even lower than those reported for $[\text{Fe}(\text{Htrz})_2(\text{trz})](\text{BF}_4)$. It is more comparable to those reported for $[\text{Fe}(\text{Htrz})_3](\text{BF}_4)_2(\text{H}_2\text{O})$.³³

To the best of our knowledge, our 40 μm -long crystals are of the largest size ever synthesized. Hence, the transition temperatures obtained in the present study, *i.e.*, $T_{1/2\uparrow} = 338$ K, $T_{1/2\downarrow} = 360$ K and $\Delta T_{1/2} = 22$ K, should serve as state-of-the-art bulk references.

2.5.1 Electrochemical analysis. The cyclic voltammogram, *i.e.*, current *versus* bias voltage shown in the left panel of Fig. 10, reveals a redox activity of a $[\text{Fe}(\text{Htrz})_2(\text{trz})](\text{BF}_4)$ thin film deposited on a FTO-coated glass. It exhibits the anodic peak at $E_{\text{pa}} \sim 0.7$ V and cathodic peak at $E_{\text{pc}} \sim 0.9$ V. The corresponding redox potential $E_{\text{redox}} = (E_{\text{pa}} + E_{\text{pc}})/2$ is ~ 0.8 V. This can be due to the reduction/oxidation of 1,2,4-triazole.⁴⁴

Fig. 10 (right) shows the electric charge accumulated in or released from the $[\text{Fe}(\text{Htrz})_2(\text{trz})](\text{BF}_4)$ thin film upon alternating the bias voltage between 0 and 1.2 V at ten second interval for the first five cycles (the red curves labelled as 1st to 5th) and the last five cycles (the blue curves labelled as 96th to 100th). Charging and discharging occurs by alternating the bias voltage between 1.2 V and 0 V at ten second interval. The maximum



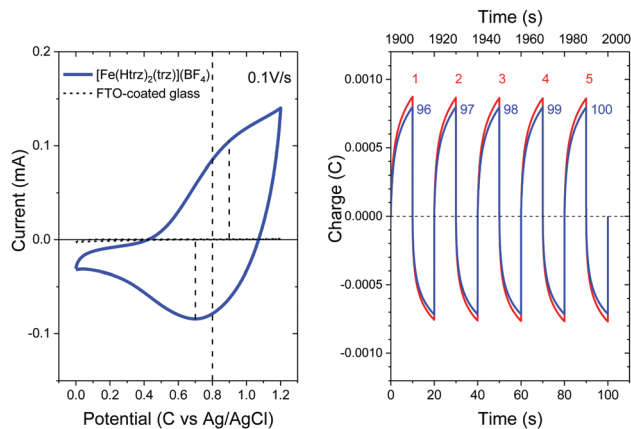


Fig. 10 (left) Cyclic voltammogram of a $[\text{Fe}(\text{Htrz})_2(\text{trz})](\text{BF}_4)$ thin film deposited on a FTO-coated glass (blue solid curve) in comparison to the reference data measured on a FTO-coated glass (black dashed curve). (right) The electric charge accumulated in or released from the $[\text{Fe}(\text{Htrz})_2(\text{trz})](\text{BF}_4)$ thin film upon alternating the bias voltage between 0 and 1.2 V at ten second interval for the first five cycles (the red curves labelled as 1st to 5th) and the last five cycles (the blue curves labelled as 96th to 100th).

charge accumulated is reduced only by $\sim 8\%$ after 100 cycles, demonstrating an excellent durability of the thin film against the oxidation–reduction reaction. Raman spectroscopy upon electrochemical doping, or spectroelectrochemistry, was also performed, but no change was observed in the Raman modes (for more details, see Fig. S2, ESI†).

2.6 Electrical conductance

Our crystals synthesised at $C = 0.3 \text{ M}$ and $T = 55^\circ \text{C}$ are of $40 \mu\text{m}$ long, which is large enough to bridge a bulky electrode gap of $30 \mu\text{m}$ (see the inset in Fig. 11) so that reliable electrical conductance measurements can be carried out. Fig. 11 shows

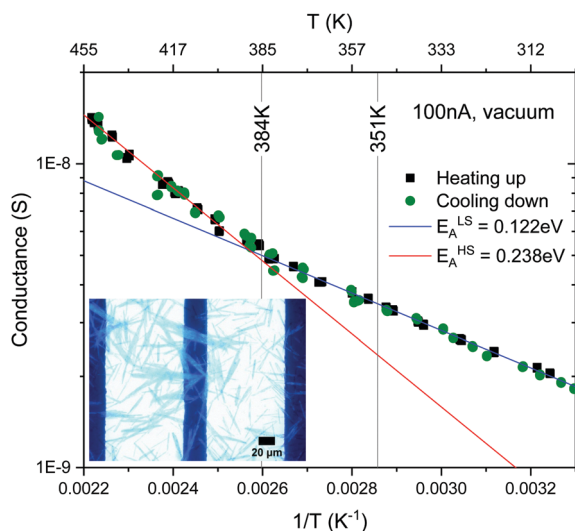


Fig. 11 Arrhenius plot of conductivity of the $[\text{Fe}(\text{Htrz})_2(\text{trz})](\text{BF}_4)$ crystalline sample plotted on a logarithmic scale as a function of temperature.

the electrical conductance plotted in the logarithmic scale as a function of temperature. The plot exhibits two linear regions with a kink at about 0.0026 K^{-1} ($\sim 384 \text{ K}$) at which the LS-to-HS transition takes place, but no discontinuity is observed upon the HS-to-LS transition at 0.00285 K^{-1} ($\sim 351 \text{ K}$). This is in contrast to previously reported DC electrical conductance data for nano- and micro-particles of $[\text{Fe}(\text{Htrz})_2(\text{trz})](\text{BF}_4)$,^{19,36–39,43,45} AC conductance⁴⁶ and DC conductance of the related complexes,⁴⁷ which exhibited thermal hystereses that coincide with respective spin thermal hysteresis. In ref. 37 and 38, sub-micrometer long rectangular crystals were aligned on interdigitated electrodes by dielectrophoresis, which allowed reliable measurements of the intrinsic electrical conductance with minimized contributions of grain boundaries. Our samples are at least 5–6 times larger than those synthesised previously so that some of the crystals directly bridge a more bulky electrode gap of $30 \mu\text{m}$. This minimises the effect of contact resistance between crystals that can be readily altered upon lattice expansion/contraction associated with spin transitions. Furthermore, if the electrical conduction is through the bulk, a much larger volume of conduction is expected in our case, which may impact on the electrical conduction mechanism.

Above and below the kink at about 0.0026 K^{-1} ($\sim 384 \text{ K}$) the data can be fit to the Arrhenius equation. The activation energies estimated for the low and high temperature domains are *ca.* $E_A^{\text{LS}} = 0.12$ and $E_A^{\text{HS}} = 0.24 \text{ eV}$, respectively. This increase by 100% in activation energy upon LS-to-HS transition is in line with the 50–70% increase reported previously on smaller crystals.¹⁹

2.7 Density functional theory

Finally, the electronic structure has been calculated based on density functional theory. Fig. 12 shows the electronic band structures of $[\text{Fe}(\text{Htrz})_2(\text{trz})](\text{BF}_4)$ in the LS state (left) and the HS state (right), respectively, along the high symmetry path Γ -Y-T-R in the Brillouin zone. In the LS state, the top of the valence bands is located at the Γ point and the bottom of the conduction bands is at the T point. The HS state has a direct band gap at the T point. The direct Kohn Sham band gap of the LS (HS) state at the T point is 2.10 eV (2.12 eV). Taking the derivative discontinuity of the exchange–correlation energy at integer-electron numbers into account, the gap energy is 2.83 eV (2.88 eV) which is greater by more than one-order of magnitude than the estimated activation energies ($E_A^{\text{LS}} = 0.12$ and $E_A^{\text{HS}} = 0.24 \text{ eV}$). The carrier injection barrier or defect levels seem to be responsible for the measured electrical conduction.

Finally, the optical band gap observed in the LS state is in line with the theoretical electronic band gap, provided that the exciton binding energy should be taken into account for a quantitative comparison (see Fig. 7). The disappearance of the absorption peak at 2.3 eV upon the LS–HS transition leads to a larger optical band gap, that is qualitatively consistent with the theoretical gap opening.



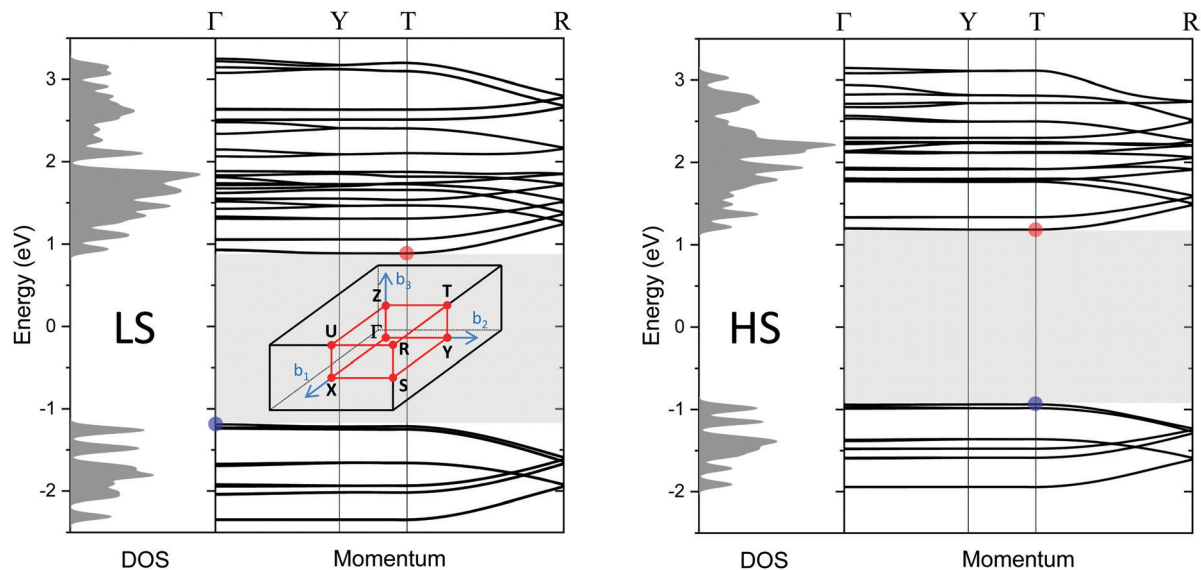


Fig. 12 Density of states and band structures for the LS (left) and HS (right) state in the momentum directions along high symmetry path Γ -Y-T-R in the Brillouin zone, calculated by using the GLLB-SC functional.^{48,49} The blue (red) solid circles are the top (bottom) of the valence (conduction) band. The shaded area represents an Kohn Sham band gap of 2.10 eV (LS)/2.12 eV (HS). Inset in the left panel: The Brillouin zone of a simple orthorhombic lattice.

3 Conclusion

The thorough systematic study on the synthesis of spin-crossover coordination polymer $[\text{Fe}(\text{Htrz})_2(\text{trz})](\text{BF}_4)$ has revealed clear dependencies of the crystal size on the solvent type, precursor concentration and the reaction temperature. The well-crystalline needles can grow as long as 80 μm by introducing a temperature gradient. It has been shown that the spin transition temperature and thermal hysteresis increase monotonically as the crystal size extends.

The *in situ* UV-vis and Raman spectroscopy measurements elucidate changes in the optical gap and vibrational modes accompanying the spin transition.

The electrochemical analysis suggests that this one-dimensional system is redox-active and stable over repeated oxidation and reduction processes. The electrical conduction across the crystals is thermally excited with activation energies of *ca.* 0.12 and 0.24 eV for the temperature range below and above 384 K, respectively. The absence of hysteresis in electrical conductance will shed light on the conduction mechanism in spin crossover systems.

The electronic band structure and the electronic gap of the $[\text{Fe}(\text{Htrz})_2(\text{trz})](\text{BF}_4)$ in the LS and HS states have been calculated based on density functional theory. The calculated electronic band gap is 2.83 eV in the LS state and 2.88 eV in the HS state. The difference between the two states is rather small as compared with the change observed in the UV-vis spectrum. Further theoretical studies with more accurate approximations are required in order to obtain good agreement between theoretical and experimental results.

The methodology for the synthesis of large crystals demonstrated in the present work provides a foundation for material sciences

centred around new and existing spin crossover materials and coordination polymers including metal-organic frameworks.

4 Methods

4.1 Synthesis

Iron(II) tetrafluoroborate hexahydrate, $\text{Fe}(\text{BF}_4)_2 \cdot 6\text{H}_2\text{O}$, purchased from Sigma-Aldrich and 1,2,4-triazole from Alfa Aesar were used as received. All solvents used were of laboratory reagent grade. The synthesis of the $[\text{Fe}(\text{Htrz})_2(\text{trz})](\text{BF}_4)$ complex was performed in two different solvents *i.e.* deionised water and methanol. The molar ratio taken for the reagents was $[\text{Fe}(\text{BF}_4)_2 \cdot 6\text{H}_2\text{O}] : [\text{triazole}] = 1 : 3$. Typically, the synthesis in methanol was performed as follows. $\text{Fe}(\text{BF}_4)_2 \cdot 6\text{H}_2\text{O}$ (0.2029 g, 0.3 M) in 2 ml of methanol was added to 1,2,4-triazole (0.1241 g, 0.9 M) in 2 ml of methanol at room temperature in a 15 ml centrifuge tube made of polypropylene. The resulting solution was maintained at room temperature for 18 hours. Then, the precipitate was separated and washed several times by decantation with fresh methanol. The same procedure was used for the synthesis in water with different precursor concentrations and temperatures. A small portion of the precipitate taken on a microscope glass slide was observed under an optical microscope. The rest was filtered off, dried and stored for further investigation. A temperature gradient was realized within a 15 ml centrifuge tube plugged in a block heater set at 45 °C. The surface of the solution in the tube was set 10 mm below the surface of the block heater. Extra-long crystals were formed right above the surface of the solution after 48 hours.

4.2 Powder X-ray diffraction

Powder X-ray diffraction measurements were carried out using a Bruker D8 Discover diffractometer with Co-K α radiation. The



crystalline sample was ground before measurements. Rietveld refinement was performed using the Full Prof Suite program.

4.3 UV-vis spectroscopy

UV-vis absorption spectra were recorded on thin films of $[\text{Fe}(\text{Htrz})_2(\text{trz})](\text{BF}_4)$ powder deposited on glass slides with a Bruker VERTEX 80v Fourier transform spectrometer.

4.4 Raman spectroscopy

Raman spectroscopy measurements were carried out on a sample placed on a hot plate with a Horiba Jobin Yvon, LabRAM HR800 with an excitation wavelength of 514.5 nm.

4.5 Magnetometry

Magnetisation measurements were carried out using a superconducting quantum interference device (SQUID) from Quantum Design. The sample after heating at 127 °C in air for 3.5 hours was placed in a gelatin capsule with the top section of the capsule being reversed to press the powder sample. The mass of the crystalline sample synthesised in water at $C = 0.3$ M and $T = 60$ °C was 15.8 mg, and that of the powder sample synthesised in methanol at $C = 0.3$ M and room temperature was 14.7 mg.

4.6 Electrochemical analysis

In our setup, silver–silver chloride (Ag/AgCl) and platinum (Pt) wires were used as the reference electrode (RE) and counter electrode (CE), respectively. The powder sample that was prepared in methanol was used to make a fine layer on a FTO-coated glass as the working electrode (WE). The electrolyte used was 0.2 M acetonitrile solution of lithium perchlorate, LiClO_4 . Cyclic voltammetry and chronoamperometry were performed using a PGSTAT 204 with Nova 2.1 software by Metrohm autolab.

4.7 Electrical conductance

Five pairs of 30 μm wide gap gold electrodes (channel width = 1 mm) were deposited on a glass plate (dimensions 25 × 18 mm × 1 mm) in a vacuum ($\sim 1 \times 10^{-6}$ mbar) by using a deposition mask (E321) purchased from Ossila. Crystals synthesised at $C = 0.3$ M and $T = 55$ °C diluted in methanol were drop cast onto the five gold electrodes. The voltage across the electrodes was measured using a Keithley 6514 system electrometer, with a constant current of 100 nA supplied by a Keithley 6221 DC and AC current source. The sample temperature was controlled by using a Lake Shore model 330 autotuning temperature controller.

4.8 Density functional theory

Structure optimisations were carried out using a Quantum ESPRESSO, an open-source plane-wave periodic density functional theory code. Norm-conserving pseudo-potentials with Perdew–Burke–Ernzerhof (PBE) exchange–correlation functional were used. Crystallographic information files, 900726 and 900727, were used to generate input files for LS and HS state calculations, respectively.¹⁶ Density functional theory calculations of the band structures were carried out by using the GPAW code which is based on the projector-augmented wave (PAW) method and the atomic simulation environment

(ASE). LCAO mode with a Monkhorst–Pack grid of (4,4,4) was used and the GLLB-SC functional was chosen in order to take the derivative discontinuity of the exchange–correlation energy at integer electron numbers into account.^{48,49} The density of states (DOS) was multiplied by normalized gaussians of 0.05 eV width.

Conflicts of interest

There are no conflicts to declare.

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Notes and references

- 1 L. S. L. Cambi, *Ber. Dtsch. Chem. Ges. A*, 1931, **64**, 2591–2598.
- 2 O. Kahn, J. Krober and C. Jay, *Adv. Mater.*, 1992, **4**, 718–728.
- 3 P. G. H. Goodwin, *Spin Crossover in Transition Metal Compounds III*, Springer, 2004.
- 4 S. Decurtins, P. Gutlich, C. Kohler, H. Spiering and A. Hauser, *Chem. Phys. Lett.*, 1984, **105**, 1–4.
- 5 S. Bonhommeau, G. Molnar, A. Galet, A. Zwick, J. Real, J. McGarvey and A. Bousseksou, *Angew. Chem., Int. Ed.*, 2005, **44**, 4069–4073.
- 6 M. A. Halcrow, *Chem. Soc. Rev.*, 2008, **37**, 278–289.
- 7 G. Molnar, V. Niel, J.-A. Real, L. Dubrovinsky, A. Bousseksou and J. J. McGarvey, *J. Phys. Chem. B*, 2003, **107**, 3149–3155.
- 8 B. Weber, W. Bauer and J. Obel, *Angew. Chem., Int. Ed.*, 2008, **47**, 10098–10101.
- 9 D.-Y. Wu, O. Sato, Y. Einaga and C.-Y. Duan, *Angew. Chem., Int. Ed.*, 2009, **48**, 1475–1478.
- 10 A. Verat, N. Ould-Moussa, E. Jeanneau, B. Le Guennic, A. Bousseksou, S. Borshch and G. Matouzenko, *Chemistry*, 2009, **15**, 10070–10082.
- 11 M. Quesada, V. A. de la Pena-O'Shea, G. Aromi, S. Geremia, C. Massera, O. Roubeau, P. Gamez and J. Reedijk, *Adv. Mater.*, 2007, **19**, 1397–1402.
- 12 M. Quesada, H. Kooijman, P. Gamez, J. S. Costa, P. J. van Koningsbruggen, P. Weinberger, M. Reissner, A. L. Spek, J. G. Haasnoot and J. Reedijk, *Dalton Trans.*, 2007, 5434–5440.
- 13 S. M. Neville, G. J. Halder, K. W. Chapman, M. B. Duriska, P. D. Southon, J. D. Cashion, J.-F. Letard, B. Moubaraki,



- K. S. Murray and C. J. Kepert, *J. Am. Chem. Soc.*, 2008, **130**, 2869–2876.
- 14 M. Quesada, F. Prins, E. Bill, H. Kooijman, P. Gamez, O. Roubeau, A. L. Spek, J. G. Haasnoot and J. Reedijk, *Chem. – Eur. J.*, 2008, **14**, 8486–8499.
- 15 V. Martinez, A. Belen Gaspar, M. Carmen Munoz, R. Ballesteros, N. Ortega-Villar, V. M. Ugalde-Saldivar, R. Moreno-Esparza and J. Antonio-Real, *Eur. J. Inorg. Chem.*, 2009, 303–310.
- 16 A. Grosjean, P. Négrier, P. Bordet, C. Etrillard, D. Mondieig, S. Pechev, E. Lebraud, J.-F. Létard and P. Guionneau, *Eur. J. Inorg. Chem.*, 2013, 796–802.
- 17 A. Michalowicz, J. Moscovici, B. Ducourant, D. Cracco and O. Kahn, *Chem. Mater.*, 1995, **7**, 1833–1842.
- 18 A. Urakawa, W. Van Beek, M. Monrabal-Capilla, J. Ramon Galan-Mascaros, L. Palin and M. Milanese, *J. Phys. Chem. C*, 2011, **115**, 1323–1329.
- 19 A. Rotaru, G. Molnár, L. Salmon, P. Demont and A. Bousseksou, *Chem. Commun.*, 2012, **48**, 4163–4165.
- 20 H. Peng, G. Molnar, L. Salmon and A. Bousseksou, *Eur. J. Inorg. Chem.*, 2015, 3336–3342.
- 21 T. Zhao, L. Cuignet, M. M. Dirtu, M. Wolff, V. Spasojevic, I. Boldog, A. Rotaru, Y. Garcia and C. Janiak, *J. Mater. Chem. C*, 2015, **3**, 7802–7812.
- 22 L. Moulet, N. Daro, C. Etrillard, J.-F. Letard, A. Grosjean and P. Guionneau, *Magnetochemistry*, 2016, **2**, 1.
- 23 A. Abrishamkar, S. Suárez-García, S. Sevim, A. Sorrenti, R. Pons, S.-X. Liu, S. Decurtins, G. Aromí, D. Aguilà, S. Pané, A. J. deMello, A. Rotaru, D. Ruiz-Molina and J. Puigmartí-Luis, *Appl. Mater. Today*, 2020, **20**, 100632.
- 24 E. Smit, B. Manoun and D. de Waal, *J. Raman Spectrosc.*, 2001, **32**, 339–344.
- 25 F. Billes, H. Endredi and G. Keresztury, *THEOCHEM*, 2000, **530**, 183–200.
- 26 J. Tuchagues, A. Bousseksou, G. Molnar, J. McGarvey and F. Varret, in *Spin Crossover in Transition Metal Compounds III*, ed. P. Gutlich and H. A. Goodwin, Topics in Current Chemistry-Series, Springer-Verlag Berlin, Heidelberg Platz 3, D-14197 Berlin, Germany, 2004, vol. 235, pp. 85–103.
- 27 N. O. Moussa, D. Ostrovskii, V. Martinez Garcia, G. Molnar, K. Tanaka, A. B. Gaspar, J. Antonio Real and A. Bousseksou, *Chem. Phys. Lett.*, 2009, **477**, 156–159.
- 28 D. Mader, S. Pillet, C. Carteret, M.-J. Stebe and J.-L. Blin, *J. Dispersion Sci. Technol.*, 2011, **32**, 1771–1779.
- 29 C. Faulmann, J. Chahine, I. Malfant, D. de Caro, B. Cormary and L. Valade, *Dalton Trans.*, 2011, **40**, 2480–2485.
- 30 Y. A. Tobon, C. Etrillard, O. Nguyen, J.-F. Létard, V. Faramarzi, J.-F. Dayen, B. Doudin, D. M. Bassani and F. Guillaume, *Eur. J. Inorg. Chem.*, 2012, 5837–5842.
- 31 G. Galle, C. Etrillard, J. Degert, F. Guillaume, J. F. Letard and E. Freysz, *Appl. Phys. Lett.*, 2013, **102**, 063302.
- 32 F. Guillaume, Y. A. Tobon, S. Bonhommeau, J.-F. Letard, L. Moulet and E. Freysz, *Chem. Phys. Lett.*, 2014, **604**, 105–109.
- 33 J. Krober, J. Audiere, R. Claude, E. Codjovi, O. Kahn, J. Haasnoot, F. Groliere, C. Jay, A. Bousseksou, J. Limares, F. Varret and A. Gonthiervassal, *Chem. Mater.*, 1994, **6**, 1404–1412.
- 34 J. Ramon Galan-Mascaros, E. Coronado, A. Forment-Aliaga, M. Monrabal-Capilla, E. Pinilla-Cienfuegos and M. Ceolin, *Inorg. Chem.*, 2010, **49**, 5706–5714.
- 35 E. Coronado, J. R. Galan-Mascaros, M. Monrabal-Capilla, J. Garcia-Martinez and P. Pardo-Ibanez, *Adv. Mater.*, 2007, **19**, 1359–1361.
- 36 C. Etrillard, V. Faramarzi, J.-F. Dayen, J.-F. Letard and B. Doudin, *Chem. Commun.*, 2011, **47**, 9663–9665.
- 37 A. Rotaru, J. Dugay, R. P. Tan, I. A. Gural'skiy, L. Salmon, P. Demont, J. Carrey, G. Molnár, M. Respaud and A. Bousseksou, *Adv. Mater.*, 2013, **25**, 1745–1749.
- 38 J. Dugay, M. Giménez-Marqués, T. Kozlova, H. W. Zandbergen, E. Coronado and H. S. van der Zant, *Adv. Mater.*, 2015, **27**, 1288–1293.
- 39 C. Lefter, R. Tan, J. Dugay, S. Tricard, G. Molnár, L. Salmon, J. Carrey, A. Rotaru and A. Bousseksou, *Phys. Chem. Chem. Phys.*, 2015, **17**, 5151–5154.
- 40 J. Manuel Herrera, S. Titos-Padilla, S. J. A. Pope, I. Berlanga, F. Zamora, J. Jose Delgado, K. V. Kamenev, X. Wang, A. Prescimone, E. K. Brechin and E. Colacio, *J. Mater. Chem. C*, 2015, **3**, 7819–7829.
- 41 M. Giménez-Marqués, M. L. García-Sanz de Larrea and E. Coronado, *J. Mater. Chem. C*, 2015, **3**, 7946–7953.
- 42 C. Bartual-Murgui, E. Natividad and O. Roubeau, *J. Mater. Chem. C*, 2015, **3**, 7916–7924.
- 43 C. Lefter, R. Tan, S. Tricard, J. Dugay, G. Molnár, L. Salmon, J. Carrey, A. Rotaru and A. Bousseksou, *Polyhedron*, 2015, **102**, 434–440.
- 44 S. Lokesh, A. Satpati and B. Sherigara, *Open Electrochem. J.*, 2010, **2**, 15–21.
- 45 F. Prins, M. Monrabal-Capilla, E. A. Osorio, E. Coronado and H. S. van der Zant, *Adv. Mater.*, 2011, **23**, 1545–1549.
- 46 C. Lefter, I. A. Gural'skiy, H. Peng, G. Molnár, L. Salmon, A. Rotaru, A. Bousseksou and P. Demont, *Phys. Status Solidi RRL*, 2014, **8**, 191–193.
- 47 C. Lefter, V. Davesne, L. Salmon, G. Molnár, P. Demont, A. Rotaru and A. Bousseksou, *Magnetochemistry*, 2016, **2**, 18.
- 48 O. Gritsenko, R. van Leeuwen, E. van Lenthe and E. J. Baerends, *Phys. Rev. A: At., Mol., Opt. Phys.*, 1995, **51**, 1944–1954.
- 49 M. Kuisma, J. Ojanen, J. Enkovaara and T. T. Rantala, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2010, **82**, 115106.

