Metal Substitution Effects on the Charge Transport and Spin Crossover Properties of [Fe$_{1-x}$Zn$_x$(Htrz)$_2$(trz)](BF$_4$) (trz = Triazole)

Constantin Lefter,‡ Simon Tricard,† Haonan Peng,‡ Gábor Molnár,‡ Lionel Salmon,‡ Philippe Demont,§ Aurelian Rotaru,∗‡ and Azzedine Bousseksou∗‡

†LCC, CNRS, and Université de Toulouse (UPS, INP), Toulouse, F-31077 cedex 4, France
‡Faculty of Electrical Engineering and Computer Science & Advanced Materials and Nanotechnology Laboratory (AMNOL), Stefan cel Mare University, 720229 Suceava, Romania
§Institut Carnot—CIRIMAT, Université de Toulouse, Toulouse, 31062 cedex 9, France

ABSTRACT: In this study we analyze the metal substitution effects on the structural, morphological, charge transport, and spin transition properties of the [Fe$_{1-x}$Zn$_x$(Htrz)$_2$(trz)](BF$_4$) (trz = triazole, x = 0, 0.26, or 0.43) compound using electron microscopy, powder X-ray diffraction, optical reflectivity, Raman, FTIR, $^{57}$Fe Mössbauer, and broadband ($10^{-2}$–$10^6$ Hz) dielectric spectroscopies. The crystal structure and the morphology of the objects remain nearly unaffected, whereas the thermal spin transition shifts from 362 to 316 K and the thermal hysteresis width decreases from 45 to 8 K for increasing values of x. For each compound the electrical conductivity drops when the iron(II) electronic configuration is switched from the low-spin to the high-spin state. A strong overall decrease in conductivity with increasing Zn concentration is also observed in both spin states. These results, together with the analysis of the charge carrier dynamics, suggest that the ferrous ions participate directly in the charge transport mechanism, explaining the strong spin-state dependence of the electrical properties in this compound.

1. INTRODUCTION

Spin crossover (SCO) complexes of 3$d^4$–$3d^7$ transition-metal ions are known to display a bistability between their high-spin (HS) and low-spin (LS) electronic configurations. The spin-state switching can be induced by various external stimuli, such as temperature, pressure, magnetic field, light irradiation, or gas sorption, and leads to a significant change of magnetic, optical, mechanical, and electrical properties. The recent progress in the nanoscale synthesis and organization of SCO compounds has opened up novel perspectives for the study of charge transport properties of these materials at the nanoscale and even at the single molecule levels. Indeed, a number of recent theoretical and experimental achievements with respect to the possible interplay between SCO and conductivity properties gives hope that in the near future SCO complexes might be integrated in functional nanoelectronic and spintronic devices.

We recently investigated the charge transport properties of the SCO compound [Fe(Htrz)$_2$(trz)](BF$_4$) and revealed a strong dependence of the dc and ac conductivities on the spin state of the iron(II) ions. The LS (1$^A_g$) state was found to be systematically more conductive than the HS (1$^T_g$) state in samples with different morphologies (nanoparticles and microrods) and sizes (ranging from 15 nm to several micrometers). In addition, the charge carrier relaxation frequencies also displayed a significant decrease when going from the LS to the HS state. We tentatively ascribed these observations to a conduction process via polaron hopping. In the frame of this model, the conductivity would rise upon the HS to LS switching (chiefly) due to an increase of the hopping frequency. This can be linked to the spin-state of the system through the variation of the phonon frequencies and/or through the variation of the hopping activation barriers. While their interpretation remains rather challenging at this stage, the accumulated experimental observations have already allowed several groups to elaborate simple nanoelectronic devices based on this compound, providing appealing perspectives for nanoscale switching and memory applications.

It is also worth noting that interesting size effects can be anticipated when going from the activated hopping to the tunneling regime because the spin transition was reported to have an opposite effect on the charge transport in these two limiting cases (typically one can expect tunneling bellow 10 nm).

The present work was motivated by the weak understanding of the physical phenomena that underlies the electrical conduction and its spin-state dependence in the title compound. The crystal structure of [Fe(Htrz)$_2$(trz)](BF$_4$) has been only recently inferred from high-quality powder X-ray diffraction (PXRD) measurements. The same orthorhombic
space group (Pnma) has been found in the two spin states. The structure consists of chains of the \([\text{Fe(Htrz)}_2(trz)]_n\) cationic motif (see Figure 1), wherein the BF₄ anions are located in cavities formed by the triazole ligands. Each chain is surrounded by six identical chains, which are connected through hydrogen bonds. An important aspect of the structure is the lack of voids, which could trap solvent molecules. The aim of the zinc(II) dilution in the present work is to break the supposed chain of conduction along the one-dimensional (1D) structure by exploiting the fact that the 3d orbitals of Zn(II) are fully occupied. On the other hand, the ionic radius of Zn(II) is close to the ionic radius of HS Fe(II), allowing for an isostructural substitution of the ferrous ions. Indeed, this type of metal dilution approach has been intensively used in spin crossover research in order to study the role of the cooperativity on the dilution approach has been intensively used in spin crossover phenomena.³¹⁻³⁴ We have studied the charge transport properties of three different powders with formulas \([\text{Fe}_{1-x}\text{Zn}_x(\text{Htrz})_2(trz)]_n[\text{BF}_4]_n\), where \(x = 0, 0.26\), or 0.43. Besides the electrical properties, each powder was carefully characterized for its morphology, crystal structure, and SCO properties.

2. EXPERIMENTAL METHODS

The undiluted microrods of the triazole coordination network \([\text{Fe(Htrz)}_2(trz)]_n(\text{BF}_4)_n\) (hereafter referred to as S1) were synthesized as described in ref 10. A 138 mg portion of 1-H,1,2,4-triazole (3 equiv) was mixed with 225 mg of Fe(BF₄)₂·6H₂O (1 equiv) in 2 mL of an HBF₄ solution at 2 wt% and left without agitation. A violet powder precipitated after 1 day. The dispersion was centrifuged and the precipitate was washed with water and dried under vacuum. The Zn-substituted samples were synthesized in a similar way, using 202 mg of Fe(BF₄)₂·6H₂O (0.9 equiv) + 23 mg of Zn(BF₄)₂·6H₂O (0.1 equiv) for sample S2 and 180 mg of Fe(BF₄)₂·6H₂O (0.8 equiv) + 45 mg of Zn(BF₄)₂·6H₂O (0.2 equiv) for sample S3. EDX (energy dispersive X-ray spectroscopy) gave Zn:Fe ratios of ca. 1:1.3 and 1:3 for S2 and S3, respectively.

The spin transition of the samples was followed through variable-temperature optical reflectivity measurements in a dry nitrogen atmosphere. The reflectance was determined by means of an optical microscope (Olympus BX51) equipped with a CCD camera (Andor IkonM) and a 543 nm band-pass filter. The sample temperature was controlled using a Linkam THMS600 liquid nitrogen cryostat. During the experiment the temperature was changed at a rate of 2 K/min.

For the structural characterization of the compounds, we used FTIR, Raman, ⁵⁷Fe Mössbauer, and PXRD methods. FTIR spectra were recorded at room temperature with a PerkinElmer Spectrum 100 spectrometer in ATR mode (resolution ca. 1 cm⁻¹) between 650 and 4000 cm⁻¹. Variable-temperature Raman spectra were acquired using a LabramHR (Horiba) Raman microspectrometer (resolution ca. 3 cm⁻¹) combined with a Linkam THMS600 cryostage. The 632.8 nm line of a He–Ne laser (0.1 mW) was focused on the sample by a x50 objective (numerical aperture = 0.5), which was also used to collect the scattered photons. These were filtered by an edge filter, but were not analyzed for their polarization state due to the random orientation of the particles in the sampled volume. Zero-field ⁵⁷Fe Mössbauer spectra were recorded using a flow-type liquid nitrogen cryostat and a conventional constant acceleration type Mössbauer spectrometer. γ-rays were provided by a ⁶⁰Co (Rh) source (ca. 20 mCi). The samples (~30 mg) were enclosed in a Plexiglas sample holder with an optimized diameter. The spectrum deconvolutions were done with the assumption of Lorentzian line shapes using the Recoil software. The PXRD patterns of the samples have been recorded at room temperature using a Panalytical MDX XpertPro powder diffractometer [Cu Kα, Ge(111) monochromator, X'Celerator detector] within the range 5°–80° by using a scan speed of 30°/h.

Particle size was determined by transmission electron microscopy (TEM) using a JEOL JEM-1011 (100 kV) instrument. TEM samples were prepared by placing a drop (10 μL) of the nanoparticles suspended in dichloromethane on a carbon-coated copper grid. High-resolution imaging was performed on a JEOL JSM 2100F (200 kV) equipped also with a SDD Bruker EDX detector (resolution 127 eV) for elemental analysis. Alternating current conductivity measurements were carried out as a function of frequency (10⁻²–10⁶ Hz) and temperature (270–400 K) by means of a BDS 4000 broadband dielectric spectrometer coupled to a Quatro Cryosystem (Novocontrol Technologies, Hundshagen, Germany) at an applied ac voltage of 1 V. Frequency sweeps were carried out isothermally. For electrical measurements, the powder samples were compacted between two stainless steel electrodes in a Teflon sample holder.

3. RESULTS AND DISCUSSION

3.1. Zn Substitution Effects on the Structure, Morphology, and Spin Transition Properties. Representative TEM images of the samples are shown in Figure 2. Each sample displays similar particle morphology, forming microrods with an average length of ca. 2 μm and a diameter of 200–300 nm. The same size and shape of the particles of the different samples is very important in order to be able to compare their conductivity. In conjunction with the TEM observations, EDX measurements were also carried out in order to evaluate the Zn/Fe ratio in the samples and to investigate the spatial distribution of the metal ions, which is also an important issue. The analysis performed on different rods for each sample revealed an average of 26 at% of Zn in S2 and 43 at% of Zn in S3, which are therefore identified as \([\text{Fe}_{0.7}\text{Zn}_{0.3}(\text{Htrz})_2(trz)]_n[\text{BF}_4]_n\) and \([\text{Fe}_{0.5}\text{Zn}_{0.5}(\text{Htrz})_2(trz)]_n[\text{BF}_4]_n\), respectively. The TEM-EDX analysis revealed also a homogeneous distribution of the Zn and Fe ions within the volume of the microrods, and no sign of ion segregation was observed (see Figure S1 in the Supporting Information).

The insertion of Zn ions within the crystal structure was investigated through X-ray diffraction and multiple spectroscopic techniques (FTIR, Raman, Mössbauer spectroscopies). The powder X-ray diffractograms (Figure 3) show a globally

![Figure 1. Schematic chain structure of \([\text{Fe(Htrz)}_2(trz)]_n\).](image-url)
similar pattern for each compound, with some differences for sample S3. For samples S1 and S2, the main diffraction peaks are grouped for values of 2θ at 10° and 11°; 18° and 19°; and 24°, 25°, and 26°. The additional diffraction peaks visible in the spectra of sample S3 are consistent with a small HS fraction at room temperature in this compound (vide infra). Indeed, as was shown in ref 10, the LS to HS transition involves a shift of the diffraction peaks toward lower 2θ angles, due to the lattice expansion. In particular, the shifts of the Bragg peaks around 24°−26° upon the SCO are rather important. This effect is clearly observed in sample S3, with the shoulders at lower angles. Overall these PXRD results can be interpreted in agreement with the EDX analysis, indicating that the Zn ions do not segregate in our compounds and instead replace some of the Fe centers to give closely isosctructural compounds.

Vibrational spectroscopy is also a useful technique to analyze the impact of metal dilution on the structure of our compounds and is a straightforward way to observe the spin transition with temperature. FTIR and Raman spectra of the three compounds are presented in the Supporting Information (Figures S2 and S3 and Tables S1 and S2). The most relevant conclusions can, however, be inferred from the temperature-dependent low-frequency Raman spectra, which are shown in Figure 4. As discussed previously, the Raman spectra of the compound [Fe(Htrz)2(trz)](BF4) exhibit characteristic changes between 100 and 300 cm⁻¹ due to the spin transition. At 293 K (LS state) the sample S1 displays intense Raman modes around 135, 197, 211, 285, and 299 cm⁻¹. When going to the HS state (413 K), one observes Raman peaks around 105, 136, 150 (shoulder), 180, and 190 cm⁻¹. These spectra are essentially the same as those reported in ref 35. The same peaks and the same changes also appear in the Zn-diluted compounds, confirming their similar structure. (See also the midfrequency spectra in Figure S3 of the Supporting Information.) While the HS spectra of the three compounds are almost identical, in the LS state spectra of the diluted compounds we observe additional peaks around 160−166 cm⁻¹ and a broadening of the peaks at 135 and 245 cm⁻¹ in both samples (S2 and S3). This difference between the HS and LS spectra can be explained by the fact that the ionic radius of Zn(II) matches that of the HS Fe(II).

The high-temperature Raman spectra are characteristic of the HS state for each sample without any significant residual LS fraction, which can be inferred clearly from the complete disappearance of the Raman peaks at 285 and 299 cm⁻¹. On the other hand, even if the low-temperature Raman spectra are clearly characteristic of the LS state, it is difficult to estimate if the transition is complete in this direction due to the spectral overlaps. To verify if there is a residual HS fraction in the samples, we acquired ⁵⁷Fe Mössbauer spectra at low temperatures (i.e., in the LS state). These spectra are shown in Figure 5 and the hyperfine parameters are collected in Table 1. In each case the spectrum can be properly fitted by only one doublet, characteristic of the LS form of Fe(II). From these Mössbauer and Raman data we can thus conclude that the spin transition of the ferrous ions is complete in both the heating and cooling modes for each sample.

This spin transition in the samples is also visible through their optical properties, since their color turns from violet to white upon heating. This bleaching of the ¹A → ¹T ligand-field absorption band (centered at ca. 545 nm) is completely reversible and provides thus a convenient way to follow the SCO phenomena in our samples. The thermal variation of the optical reflectance in the green spectral region is shown in Figure 6 for each sample. As expected, the spin transition is shifted toward lower temperatures, and the hysteresis width decreases for increasing Zn substitution (Table 2). Such metal dilution effects are well-known and widely used by the spin crossover community to tune and investigate spin transition properties (see, for example, refs 1, 2, and 32−35). A short
report on the Zn dilution of the title compound, wherein a similar—though somewhat less pronounced—shift of the spin transition was previously reported for a 20% Zn doping.\textsuperscript{36} The downshift of the spin transition upon Zn doping can be explained by the fact that the ionic radius of the Zn\textsuperscript{2+} ion (96 pm) is much closer to the ionic radius of Fe\textsuperscript{2+} ions in the HS state (92 pm) than in the LS state (75 pm);\textsuperscript{37} i.e., the LS lattice is destabilized due to the misfit of the Zn ions. On the other hand, the hysteresis in SCO compounds is known to occur due to the elastic interactions between the SCO centers. Since the Zn dilution leads to a decrease of the concentration of the SCO-active iron ions, the cooperativity of the system decreases, which is reflected by the shrinking of the hysteresis as well as by an increasingly gradual transition.\textsuperscript{1,2,32−38}

3.2. Zn Substitution Effects on Charge Transport and Dielectric Properties.

Broadband dielectric spectroscopy provides access to a wide range of electrical properties. In particular, the complex conductivity $\sigma^*(\omega,T)$ and its frequency and temperature dependences give important information about the charge transport mechanism. The frequency dependence of the real part, $\sigma'\!(\omega)$, of the complex conductivity is reported in Figure 7a−c for each compound at different temperatures both in the heating and cooling branches. The shape of the dispersion curves and the sample conductivity are strongly dependent on the spin state and the Zn dilution. For S1 the frequency dispersion of the electrical conductivity is strongly dependent on the spin state and temperature (Figure 7a), in good agreement with our previous reports on the pure complex.\textsuperscript{10,11} The conductivity $\sigma'\!(\omega)$ shows, at all temperatures, a typical low-frequency plateau, characteristic of a nonlocalized or a long-range charge carrier mobility. Above the crossover frequency, $\omega_c$, the conductivity $\sigma'\!(\omega)$ increases with increasing frequency, obeying a power law. In other words, the ac conductivity spectra follow Jonscher's power law\textsuperscript{39}

$$\sigma'\!(\omega) = \sigma_{dc} + A\omega^n = \sigma_{dc} \left[1 + \left(\frac{\omega}{\omega_c}\right)^n\right]$$  

(1)
where $\sigma_{dc}$ is the dc conductivity; $\omega_c$ is the crossover frequency from the dc to the dispersive region, representing the hopping frequency of charge carriers; $A$ is a temperature- and frequency-dependent parameter; and $n$ is an exponent dependent on both frequency and temperature in the range $0 < n \leq 1$. This behavior is characteristic of the charge transport in disordered materials and was interpreted by Jonscher as a universal dynamic response (UDR). For S2 and S3, the plateau at low frequency is strongly reduced and the dispersion of the ac conductivity shifts to lower frequency, i.e., $\omega_c$ decreases (Figure 7b,c), in agreement with the more insulating character of the diluted samples. Indeed, the conductivity $\sigma'$ of the samples at $10^{-2}$ Hz (dc regime) and at 293 K (LS state) is ca. $10^{-10}$ S cm$^{-1}$ for S1, $10^{-13}$ S cm$^{-1}$ for S2, and $10^{-16}$ S cm$^{-1}$ for S3. This huge decrease in conductivity upon Zn substitution is observed in the whole investigated frequency and temperature range, i.e., in both spin states and conductivity regimes (ac and dc).

The thermal dependence of the real part of conductivity at different values of $E_f$ for samples S1–S3 at the LS to the HS state, but no clear correlation exists with the Zn dilution; at room temperature (LS state) the values of $\omega_c$ increases upon increasing the percentage of the Zn dilution. This observation is in good agreement with the decrease of $E_f$ obtained from the Arrhenius fit in the LS and HS states at different frequencies are reported in Figure 9.

Even if the quality of the fit becomes rather low for samples S2 and S3, we can draw several important conclusions. First, the activation energy in the LS state is systematically lower than in the HS state. Second, the Zn dilution does not influence considerably the activation barriers of the conductivity. The activation energy has no frequency dependence below the cutoff frequency ($\omega_c$), but it decreases strongly for higher frequencies in each sample and in both spin states. The frequency at which the activation energy varies decreases upon increasing the percentage of the Zn dilution. This observation is in agreement with the decrease of $\omega_c$ with the dilution.

We have also extracted the values of $\omega_c$ and $n$ by a nonlinear fitting procedure using eq 1 for each sample and each temperature. The temperature dependence of these two parameters are reported in Figures S5 and S6 in the Supporting Information. In these figures we observe the same hysteretic behavior related to the spin transition phenomenon. There is a clear decrease of the power law exponent $n$ when going from the LS to the HS state, but no clear correlation exists with the Zn dilution; at room temperature (LS state) the values of $n$ fall between 0.5 and 0.6 for each sample. The thermal variation of the hopping frequency $\omega_c$ brings more information about the charge transport mechanism in the samples. Most importantly, the hopping frequency decreases very significantly with the Zn dilution. In the LS state we found the hopping frequency to be $138$ s$^{-1}$ for S1, $0.12$ s$^{-1}$ for S2, and $0.01$ s$^{-1}$ for S3. Furthermore, $\omega_c$ is always higher in the LS state, but its variation is decreasing with increasing amount of Zn impurities. Overall, the behavior of $\omega_c$ shows a clear correlation with that

$$\sigma' = \sigma_0' \exp\left(-\frac{E_f}{k_B T}\right)$$

Figure 8. Temperature dependence of the ac conductivity for samples S1–S3 at low (a, c, e) and high frequencies (b, d, f).

Figure 9. Frequency dependence of the conductivity activation energy for samples S1–S3 in the two spin states.
of the conductivity $\sigma'$. We can thus suggest that the strong decrease of the conductivity with the Zn dilution as well as by the LS to HS transition is primarily governed by the decrease of the hopping frequency.

The dispersion behavior of the electrical conductivity in the frequency domain can also be interpreted in terms of conductivity relaxation time using the electric modulus, $M^* = 1/e^{\omega\tau}$, representation.\footnote{40,41} \begin{equation} \text{M}^* = M' + iM'' = M_\infty - \frac{(M_\infty - M_0)}{[1 + (i\omega\tau_M)^\alpha \beta]} \end{equation} where $M_\infty = 1/\epsilon_\infty$ and $M_0 = 1/\epsilon_0$ are the limiting low- and high-frequency moduli, $\tau_M$ is a mean characteristic relaxation time related to the peak frequency $\omega_M$, while $\alpha$ and $\beta$ are fractional shape parameters ($0 < \alpha, \beta < 1$). The inverse of the maximum peak frequency of the $M^*$ spectra represents the time scale of the transition from the long-range to short-range mobility and is defined as the characteristic conduction relaxation time ($\tau_M = 1/\omega_M$). The temperature dependence of the relaxation peak frequencies is shown in Figure 10. For each sample, we can observe the hysteresis associated with the spin transition due to a decrease of $\omega_M$ when going from the LS to the HS state, even if this variation is attenuated for increasing Zn doping level. One can note also that the relaxation frequencies for the same temperature and same spin state decrease drastically when Zn impurities are inserted. In the LS state the values of $\omega_M$ and $\omega_\infty$ are $5 \times 10^3$ s$^{-1}$ and $4 \times 10^4$ s$^{-1}$ for sample S1, $4.5 \times 10^3$ s$^{-1}$ and $2.7 \times 10^3$ s$^{-1}$ for sample S2, and $8 \times 10^2$ s$^{-1}$ and $2.3 \times 10^2$ s$^{-1}$ for sample S3.

It is tempting to correlate in a more quantitative manner the thermal behavior of $\omega_M, \omega_\infty$, and $\omega_\sigma$, which are all thermally activated and show qualitatively similar evolution both with the spin transition and Zn dilution. Indeed, the temperature dependence of both $\omega_M$ and $\omega_\sigma$ obeys an Arrhenius law that can be described by the following equations:

\begin{equation} \omega_\infty = \omega_{0\infty} \exp\left(-\frac{E_h}{k_B T}\right) \end{equation}

\begin{equation} E_h = \omega_{0M} \exp\left(-\frac{E_M}{k_B T}\right) \end{equation}

involving $E_\sigma, \omega_{0h}$ and $E_M, \omega_{0M}$ as activation parameters, respectively. We could obtain satisfactory fits for all kinds of activation parameters only for sample S1 (Table 3). For this compound, the activation energies extracted from electric modulus analysis agree with the values of dc conductivity and hopping frequency activation energies. In other words, similar energy barriers are involved both in the relaxation and conduction processes and it can therefore be assumed that these different representations of charge transport are equivalent. This conclusion is also supported by the very similar values obtained for the frequency pre-exponential factors $\omega_{0\sigma}, \omega_{0MT}$, and $\omega_{0MT}$. When comparing the spin-state dependence of activation parameters there is an obvious decrease of the activation energy when going from the HS to the LS state, while the spin-state dependence of the pre-exponential factors is less obvious. On the other hand, the activation energies (in a given spin state) are very similar in the pure and Zn-diluted samples (see Figure 9). The dramatic

![Figure 10. Frequency dependence of the loss modulus in the LS and HS states for samples S1 (a), S2 (b), and S3 (c). Temperature dependence of the two deconvoluted relaxation peak frequencies for samples S1 (d), S2 (e), and S3 (f).](image)

Table 3. Activation Parameters for Sample S1 in the HS and LS States Obtained from Arrhenius Dependence of the Direct Current Conductivity, the Hopping Frequency ($\omega_\sigma$), and the Loss Peak Frequency ($\omega_M$) in the Heating (↑) and Cooling (↓) Modes

<table>
<thead>
<tr>
<th>spin state</th>
<th>$E_\sigma$ (eV)</th>
<th>$\sigma_0$ (S cm$^{-1}$)</th>
<th>$E_\omega$ (eV)</th>
<th>$\omega_{0\infty}$ (s$^{-1}$)</th>
<th>$E_{0M}$ (eV)</th>
<th>$\omega_{0MT}$ (s$^{-1}$)</th>
<th>$E_{0MT}$ (eV)</th>
<th>$\omega_{0MTD}$ (s$^{-1}$)</th>
</tr>
</thead>
<tbody>
<tr>
<td>HS</td>
<td>0.61↑</td>
<td>$1 \times 10^{-2}$</td>
<td>0.59↑</td>
<td>$2 \times 10^{10}$</td>
<td>0.67↑</td>
<td>$2 \times 10^{11}$</td>
<td>0.59↑</td>
<td>$2 \times 10^{10}$</td>
</tr>
<tr>
<td></td>
<td>0.61↓</td>
<td>$1 \times 10^{-2}$</td>
<td>0.58↓</td>
<td>$1 \times 10^{10}$</td>
<td>0.64↓</td>
<td>$3 \times 10^{10}$</td>
<td>0.58↓</td>
<td>$1 \times 10^{10}$</td>
</tr>
<tr>
<td>LS</td>
<td>0.51↑</td>
<td>$6 \times 10^{-2}$</td>
<td>0.56↑</td>
<td>$3 \times 10^{11}$</td>
<td>0.47↑</td>
<td>$4 \times 10^{10}$</td>
<td>0.57↑</td>
<td>$5 \times 10^{11}$</td>
</tr>
<tr>
<td></td>
<td>0.48↓</td>
<td>$1 \times 10^{-2}$</td>
<td>0.53↓</td>
<td>$1 \times 10^{11}$</td>
<td>0.45↓</td>
<td>$7 \times 10^{9}$</td>
<td>0.53↓</td>
<td>$1 \times 10^{11}$</td>
</tr>
</tbody>
</table>
decrease of $\sigma_{dc}$, $\sigma_{ac}$, and $\omega_c$, with the dilution cannot be thus related to a change of the activation barriers. We suggest, therefore, that these changes are related either to a decrease of the charge carrier density (number of iron centers) and/or to a change of the hopping distance and the associated active phonon modes.

4. CONCLUSIONS

Electrical properties of a Zn-diluted series of the [Fe$_{x-1}$Zn$_x$(Htrz)$_2$(trz)]($\text{BF}_4$) spin crossover compound were studied by broadband dielectric spectroscopy in a large frequency and temperature range. This study represents the first investigation of the influence of substitution of the "active" Fe centers by "inactive" Zn ions on the charge transport properties of a spin crossover compound. The metal substitution was found to be homogeneous, and the obtained compounds appeared closely isostuctural. The iron ions kept their spin transition properties in the diluted samples, but as expected, a loss of cooperativity and a shift of the spin transition toward lower temperatures were observed with increasing the iron dilution by zinc. We have demonstrated that spin crossover phenomena can be detected through the temperature dependence of ac conductivity and dielectric loss, and we show that virtually all material-dependent electrical parameters, such as electrical conductivity, electric modulus, cutoff frequency, relaxation peak frequency, display a spin-state dependence. In particular, we have shown that the spin transition from the LS to the HS state led to a systematic decrease of the electrical conductivities and carrier hopping frequencies, which were related to the higher values of the activation energy in the HS state. The Zn substitution of active iron centers does not change the activation barriers but leads to an important decrease of the charge carrier hopping frequencies, which is reflected by the drop of the electrical conductivity by ca. 6 orders of magnitude (for $x = 0.43$). Taking into account the close structural and morphological similarity of the pure and Zn-diluted samples, these results indicate that the ferrous ions with open 3d$^6$ electronic shell directly participate in the charge transport process, in contrast to the Zn(II) ions, which have a closed 3d$^{10}$ shell.

■ ASSOCIATED CONTENT

Supporting Information
EDX, FTIR, and Raman data and additional analysis of the conductivity measurements. This material is available free of charge via the Internet at http://pubs.acs.org.

■ AUTHOR INFORMATION

Corresponding Authors
*A.R. e-mail: rotaru@eed.usv.ro.
*A.B. e-mail: azeddine.bousseksou@lcc-toulouse.fr.

Notes
The authors declare no competing financial interest.

■ ACKNOWLEDGMENTS

This work was founded by the joint French–Romanian project ANR-UEFISCDI, contract nos. 9RO-FR/01.02.2013 and ANR-12-IS07-0003-01. S.T. thanks the European Community for financial support (Grant PCIG11-GA-2012-317692). C.L. thanks the French Ministry of Foreign Affairs (MAEE) for an Eifel Scholarship.

■ REFERENCES

(14) Etrillard, C.; Faramarzi, V.; Dayen, J.-F.; Letard, J.-F.; Doudin, B. Photocoduction in Fe(Htrz)$_2$(trz)($\text{BF}_4$)$_2$H$_2$O Nanocrystals. Chem. Commun. 2011, 47, 9663–9665.
(20) Meded, V.; Bagrets, A.; Fink, K.; Chandrasekar, R.; Ruben, M.; Evers, F.; Bernard-Mantel, A.; Seldenthuis, J. S.; Beukman, A.; van der Zant, H. S. J. Electrical Control over the Fe(II) Spin Crossover in a
Pechev, S.; Lebraud, E.; Le and Spin Crossover in the Polymeric Material [Fe(Htrz)2(trz)](BF4) Low-Spin Conversion in the Molecular System [FeIII(qsal)2][Ni-
Sato, O. Electrical Conductivity Modulation Coupled to a High-Spin-
2154
Low-Spin 1A1 Codjovi, E. Study of Impurities Effect in Spin Crossover Compounds
of Molecular Spin-Crossover Complex Fe(phen)2(NCS)2 Thin Films.
Boukari, S.; Bowen, M.; Kemp, N. T.; Viart, N.; Rogez, G.; et al. Study
Beads of Iron(II) Complexes.
Student, K.; Grohmann, A. Spin-State Patterns in Surface-Grafted
Complexes with Tetrathiafulvalene Moieties.

Crossover Behavior and Electrical Conduction Property in Iron(II)

Using First Order Reversal Curves (FORC) Method. Polyhedron 2007,
26, 1820–1824.

(38) Rotaru, A; Durtu, M. M.; Enachescu, C.; Tanasa, R; Linares, J;
Stancu, A; Garcia, Y. Calorimetric Measurements of Diluted Spin
Crossover Complexes [FeMII(br)3(NCS)2]·H2O with MII = Zn and

267, 673–679.

(40) Macedo, P. B.; Moynihan, C. T.; Bose, R. Role of Ionic

(41) Gerhardt, R. Impedance and Dielectric Spectroscopy Revisited:
Distinguishing Localized Relaxation from Long-Range Conductivity. J.

(42) Molnar, G.; Cobos, S.; Mahfoud, T.; Vertelman, E. J. M.; van
Koningsbruggen, P. J.; Demont, P.; Boussesou, A. Interplay Between
the Charge Transport Phenomena and the Charge-Transfer Phase
Transition in RbMn[Fe(CN)6]·H2O. J. Phys. Chem. C 2009, 113,
2586–2593.

(43) Rouahi, A; Kahouli, A.; Challali, A.; Beslaid, M. P.; Vallee, C.;
Yangui, B; Salimy, S.; Goullet, A.; Sylvestre, A. Impedance and Electric
Modulus Study of Amorphous TiTaO Thin Films: Highlight of the

(44) Havrilak, S.; Negami, S. A Complex Plane Representation of
Dielectric and Mechanical Relaxation Processes in some Polymers.