**Powder Sample Susceptibility for Single Ion Magnets with \( S = 1, 3/2 \) with Rhombic Anisotropy**

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In this paper, the general analysis of powder sample magnetic susceptibility of single ion magnets displaying axial (\( D \)) and rhombic (\( E \)) local anisotropy, and characterized by the isotropic spectroscopic tensor \( g \), is conducted. The aim is to obtain exact expressions for the temperature dependence of magnetic susceptibility to extract both anisotropy parameters \( D \) and \( E \) from the powder susceptibility data. Two cases are considered, with integer (\( S = 1 \)) and half-integer (\( S = 3/2 \)) spin. In the former, the procedure based on the low-temperature behavior of the susceptibility signal, which effectively reduces the number of fitting parameters, is proposed. By contrast, in the case of the SIM with \( S = 3/2 \), it is explicitly demonstrated that the powder susceptibility depends on the combined anisotropy parameter \( \Delta = \sqrt{D^2 + 3E^2} \), precluding the full resolution of the axial and rhombic anisotropy constants.

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1. Introduction

New materials are recently being sought for further development of data storage density. One of the most promising candidates are single ion magnets (SIM) that theoretically could be used to encode binary information in a single molecule. Concept of magnetic data storage requires bistable remnant magnetization with an effective energy barrier when switching between \( +z \) and \( -z \) states of the magnetic moment \([1]\). The origin and size of an energy barrier directly lie in magnetic anisotropy, a phenomenon in which a molecule is easier to magnetize along a certain crystallographic axis. Basically, the stronger is the magnetic anisotropy, the higher is the energy barrier.

SIM’s are mononuclear complexes that possess sufficient magnetic anisotropy and spin to retain magnetization for a specific time in the low-temperature range \([2]\). They comprise of a single magnetic ion whose properties are modified by the surrounding ligand field causing magnetic anisotropy. In the absence of external magnetic field, this phenomenon is called zero field splitting (ZFS), which is present only when the spin ground state of a molecule is greater than 1/2 and if the symmetry is lower than cubic \([3]\). The relevant Hamiltonian \( \hat{H} \) for an isolated magnetic moment in an anisotropic environment has the following form:

\[
\hat{H} = D \left( \hat{S}_z^2 - \frac{1}{3}S(S+1) \right) + E \left( \hat{S}_x^2 - \hat{S}_y^2 \right),
\]

where \( D \) and \( E \) are the axial and rhombic ZFS parameters respectively, \( S \) is the spin quantum number, and \( \hat{S} = (\hat{S}_x, \hat{S}_y, \hat{S}_z) \) denotes the spin operator components.

For the integer spin system, ZFS completely removes the degeneracy of the \( S \) multiplet if the terms \( D \) and \( E \) are non-zero. In the case of the half-integer spins, according to the Kramers theorem \([4]\), ZFS splits \( 2S + 1 \) energy levels into degenerate pairs with the same intrinsic angular momentum \( |M_S| \). Degeneracy can be further removed only by the magnetic field.

The development in the field of coordination compounds displaying considerable magnetic anisotropies requires appropriate theoretical models. The approach presented in \([5]\) allows deriving the exact formulae for the full magnetic susceptibility tensor with both ZFS parameters and an arbitrary spectroscopic tensor \( g \) for spin models with \( S = 1, 3/2, \) and \( 5/2 \). In this paper, the special cases of powder sample susceptibility with an isotropic tensor \( g \) for the integer \( S = 1 \) and the half-integer \( S = 3/2 \) spins are discussed in detail. The aim is to allow one to obtain both ZFS parameters from the experimental powder data with simplified formulae and reduced number of fitting parameters.

2. Results

The temperature dependence of the zero-field powder sample magnetic susceptibility in the system with the rhombic anisotropy is presented in this work. The problem is divided into two parts, one representing an integer spin system (\( S = 1 \)), and the other one, a half-integer spin system (\( S = 3/2 \)). The susceptibility of a powder sample was obtained from the corresponding formulae taken from \([5]\) by appropriately averaging over the continuum of single crystal orientations yielding

\[
\chi(T) = \frac{1}{3} \text{Tr} \left[ \hat{\chi}(T) \right] = \frac{1}{3} \left( \chi_{xx}(T) + \chi_{yy}(T) + \chi_{zz}(T) \right).
\]

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2.1. The case of $S = 1$

If molar quantities are considered, the isotropic magnetic susceptibility $\chi_{\text{iso}}(T)$ of an isolated magnetic moment reads

$$\chi_{\text{iso}}(T) = N_A\mu_B^2 g^2 S(S + 1)/\beta,$$

where $N_A$ denotes the Avogadro constant, $\mu_B$ is the Bohr magneton, $g$ is the Landé factor, $k_B$ is the Boltzmann constant, and $\beta = 1/(k_B T)$ is the thermodynamic beta. Assuming $g = 2$ and $S = 1$, $\chi_{\text{iso}}(T)$ has a constant value of about $1 \text{ cm}^3/(\text{K mol})$. Let us introduce the parameter $\lambda = E/D$ which measures the ratio of the ZFS parameters and is limited to be $|\lambda| < \frac{1}{2}$ to ensure that $z$-axis is always either the easy or the hard axis [2]. The powder sample susceptibility with isotropic $g = \text{diag}[g, g, g]$ then reads

$$\bar{\chi}(T) = \chi_{\text{iso}}(T) F_1(\beta D, \lambda),$$

where

$$F_1(u, \lambda) = \frac{1}{u} e^u + 2 \cosh(\lambda u) \times \left[ \frac{2}{1 - \lambda^2} \frac{e^u - \cosh(\lambda u)}{u} + \frac{1 - 3\lambda^2 \sinh(\lambda u)}{1 - \lambda^2 \lambda u} \right].$$

The dimensionless quantity $u$ is defined as $u = \beta D$. In the limit of high temperature $T \to \infty$ or isotropic environment $D \to 0$ (i.e., $u \to 0$) function $F_1(\beta D, \lambda) = 1$, as expected for an isotropic magnetic susceptibility $\chi_{\text{iso}}(T)$. Figures 1 and 2 show the dependence of the powder sample susceptibility for $S = 1$ and $g = 2$ for various values of parameters $D$ and $\lambda$. When $D > 0$, it is hard to distinguish between curves with the same $D$ and different $\lambda$, which effectively precludes the estimation of $E$. On the other hand, when $D < 0$ and is sizable enough, all the curves are well separated, as shown in Fig. 2.

![Fig. 1. Temperature dependence of the powder sample susceptibility $\bar{\chi}(T)$ given in (4), for spin value $S = 1$ with $\lambda = 0.01$ (black), $\lambda = 0.1$ (red), and $\lambda = 0.3$ (blue), and the various positive axial anisotropy parameters $D > 0$.](image1)

![Fig. 2. As in Fig. 1, but for the various negative axial anisotropy parameters $D < 0$.](image2)

One can extract the leading term proportional to $1/u$ as the temperature approaches zero (i.e., $u \to \pm \infty$). Two distinct cases corresponding to the different sign of parameter $D$ must be considered separately. For $D < 0$ the asymptotic formula reads

$$F_1(-\beta |D|, \lambda) \propto \frac{3 |\lambda| + 1}{2|\lambda|(1 + |\lambda|)} \frac{1}{|D|},$$

while for $D > 0$

$$F_1(\beta D, \lambda) \propto \frac{2}{1 - \lambda^2} \frac{1}{\beta D}.$$

Combining Eq. (3) with either (6) or (7) indicates that the limiting value of $\bar{\chi}(T \to 0)$ is a non-zero constant depending on the ZFS parameter $\lambda$. Thus, the product $T \cdot \bar{\chi}(T)$ displays a linear temperature dependence in this limit. Let us define a quantity that can be estimated directly from the powder sample susceptibility data

$$A = \frac{\partial}{\partial T} \left. \frac{T \cdot \bar{\chi}(T)}{T = 0} \right|,$$

which is a slope of the $T \cdot \bar{\chi}(T)$ curve in the zero temperature limit. With this new variable, the ZFS parameter ratio $\lambda$ can be expressed in terms of $D$, $A$, and $g$:

$$|\lambda| = \frac{1}{2} \left[ \frac{N_A\mu_B^2 g^2}{|D|A} - 1 \right] + \sqrt{\left( \frac{N_A\mu_B^2 g^2}{|D|A} - 1 \right)^2 + \frac{4 N_A\mu_B^2 g^2}{3 |D|A}},$$

for $D < 0$, and

$$|\lambda| = \sqrt{1 - \frac{4 N_A\mu_B^2 g^2}{3 |D|A}},$$

for $D > 0$. Note that in the above expressions $A$ is defined in units of $\text{cm}^3/\text{mol}$. This procedure reduces the number of fitting parameters of function $T \cdot \bar{\chi}(T)$ to two, i.e., the axial parameter $D$ and the spectroscopic factor $g$, provided that one has estimated the slope $A$. Consequently, the rhombic parameter $E$ can be calculated...
instantly as $E = \lambda D$. Table I shows the upper temperature limit at which the above linear approximation is still valid for certain values of parameters $\lambda$ and $D$. For $D < 0$, the larger magnetic anisotropy is (and thus $D$ and $E$), the higher is the temperature at which one can use (9) to reduce the number of fitting parameters. In the case of $D > 0$, the upper temperature limit changes only with $\lambda$ and has no substantial dependence on $D$.

We have made an attempt to check the above procedure out on the real data. We considered two transition-metal complexes $[\text{M}^{II}(\text{N}_{3}\text{N}^{2+}\text{Li}^{+} \text{THF})] \ (\text{M}=\text{Co}, \text{Ni})$ [6]. The transition metal ions Co$^{II}$ and Ni$^{II}$ in these compounds carry the spin of $S = 3/2$ and $S = 1$, respectively. The constraint fitting of the Ni congener according to the above procedure yielded only a slightly lower root mean square deviation of $1.7 \times 10^{-2}$ cm$^3$/K mol in the $D < 0$ case than in the $D > 0$ case. The low-temperature slope of the $T \cdot \chi(T)$ curve was found to amount to $A = 0.072(2)$ cm$^3$/mol. The linear low-temperature approximation together with the best fit curve are shown in Fig. 3. The best-fit parameters amount to $g = 1.979(3)$, $D = -18.5(3)$ cm$^{-1}$, and $|E| = 7(2)$ cm$^{-1}$. While the value of $E$ compares well with the value of 6(1) cm$^{-1}$ reported in [6], that of the $D$ parameter is considerably underestimated (in [6] $D = -26(1)$ cm$^{-1}$ is reported). At the same time the unconstrained fitting with all three parameters $g$, $D$, and $E$ relaxed, gives a much better root mean square deviation of $8.7 \times 10^{-3}$ cm$^3$/K mol and the parameter set, $g = 1.987(2)$, $D = -24.2(7)$ cm$^{-1}$, $|E| = 5.7(3)$ cm$^{-1}$, in a better agreement with the values reported in [6] and obtained by the simultaneous fitting of the susceptibility and magnetization data. This implies that the proposed procedure might not be a robust one, depending crucially on the precise determination of the slope $A$, which can be thwarted by the presence of the intermolecular interactions.

### 2.2. The case of $S = 3/2$

As mentioned in the first section, crystal field (electrostatic interactions with neighboring valence electrons) cannot fully remove the degeneracy for systems with the half integer spin. The resulting Kramers doublets can be further split only by applying an external magnetic field. As a consequence, even in the limit of zero magnetic field the susceptibility tensor depends on the direction of the sampling field as demonstrated in [5]. It was not the case for $S = 1$, and it is actually a characteristic feature for half-integer spins. However, by averaging over all possible directions of the sampling field one arrives at the common result comprising the arithmetic mean of the susceptibility values taken for the three orthogonal sampling field directions as in (2). The final formula for the powder sample susceptibility of the $S = 3/2$ system reads

\[
\bar{\chi}(T) = \chi_{\text{iso}}(T) F_{3/2}(\beta \Delta)
\]

with $\Delta = \sqrt{D^2 + 3E^2}$, $\chi_{\text{iso}}(T)$ is a constant with value of about 1.875 cm$^3$/K mol ($g = 2.0$) according to (3), and

\[
F_{3/2}(\beta \Delta) = \frac{1}{5} \left( 3 + 2\frac{\tanh(\beta \Delta)}{\beta \Delta} \right).
\]
In the high-temperature limit $F_{3/2}(\beta \Delta) = 1$ and in the low-temperature limit $F_{3/2}(\beta \Delta) = 3/5$, as shown in Fig. 4, above equation may look much simpler compared to the case of $S = 1$. However, since $\Delta$ combines both ZFS parameters, it is impossible to unambiguously extract both of them from a powder experiment. In fact, only $\Delta$ is attainable which gives information about the size of the magnetic anisotropy but without the distinction between the axial and rhombic ZFS parameters. In Fig. 3 the best-fit curve is shown for the Co congener. The resulting parameter $\Delta$ amounts to $23.9(7)$ cm$^{-1}$, which is lower than the value of $28(1)$ cm$^{-1}$ implied in [6]. This indicates that fitting of the powder susceptibility alone may not be fully competitive to the simultaneous analysis of the susceptibility and magnetization data.

3. Conclusions

Exact formulae for the powder sample magnetic susceptibility have been presented and discussed for SIM with integer ($S = 1$) and half-integer ($S = 3/2$) spins. In the case of $S = 1$, it was shown that both ZFS parameters, $D$ and $E$, might be distinctly obtained for $D < 0$ when magnetic anisotropy is large enough. Furthermore, the procedure, albeit not always robust, based on the linear behavior of the $T \chi(T)$ in the low temperature limit and reduced number of fitting parameters, was proposed with the discussion of conditions of applicability of such an approach. A similar approach could not be done for $S = 3/2$ because $T \cdot \chi(T)$ depends on the combination of ZFS parameters that does not allow for extracting the exact values of both $D$ and $E$ anisotropy constants simultaneously.

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