Determination of the single-ion anisotropy energy in a $S = \frac{5}{2}$ kagome antiferromagnet using x-ray absorption spectroscopy


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We report x-ray absorption and x-ray linear dichroism measurements at the Fe $L_2,3$ edges of the geometrically frustrated systems of potassium and hydronium iron jarosite. Comparison with simulated spectra, involving ligand-field multiplet calculations modeling the 3$d^2$-2p hybridization between the iron and the oxygen ligands, has yielded accurate estimates for the ligand metal-ion hybridization and the resulting single-ion crystal-field anisotropy energy. Using this method we provide an experimentally verified scenario for the appearance of a single-ion anisotropy in this nominally high-spin 3$d^5$ orbital singlet $^5S$ system, which accounts for features of the spin-wave dispersion in the long-range-ordered ground state of potassium iron jarosite.

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I. INTRODUCTION

The jarosite group of minerals has received much attention for being close to ideal realizations of the kagome antiferromagnet. The kagome topology, which is named after a Japanese basket weaving pattern, frustrates the antiferromagnetic Néel ordering.1 Due to the resulting “underconstraint” neither in the quantum case of $S = 1/2$ nor in the classical limit is the kagome antiferromagnet expected to show a symmetry-breaking transition, even at $T = 0$ K.2–4 Physical realizations of this system are highly valued because they allow for the investigation of the Mott insulating phase in the absence of Néel-type magnetic order. The jarosite group of general stoichiometry $AM_n(SO_4)_2(OH)_m$, where $A$ can be—among others—K, Na, Rb, Ag, NH$_3$, or H$_3$O, provides physical models for the kagome antiferromagnet with spin $S$ = 3/2 and 5/2, where the $M$ site is occupied by trivalent Cr (Refs. 5–7) and Fe,5,8–14 respectively. $M = V$ is a $S = 1$ system with ferromagnetic near-neighbor interactions.15,16 Figure 1 illustrates the jarosite structure of well-separated kagome layers consisting of $MO_6$ octahedra. Potassium iron jarosite is representative for most of the iron analogs. In this system a transition to a noncollinear long-range-ordered state is observed at 64 K despite that the magnitude of the Curie-Weiss temperature $\Theta_{CW}$ is as high as $\approx 800$ K.11,17 The ground state has shown to be a so-called $q = 0$ state of positive chirality, in which the spins lie within the kagome plane and all point either in or out of each shared kagome triangle.9,12,13,18,19 The large figure $\Theta_{CW}/T_N = 12.5$ is an indication of relatively strong geometric frustration,20 but clearly the observed long-range-ordered ground state points to additional terms in the Hamiltonian of this kagome antiferromagnet beyond near-neighbor exchange. It has been shown that the $q = 0$ ground state can arise due to magnetic anisotropies such as easy-plane single-ion crystal-field (CF) anisotropy $D_{CF}S_z^2 - E_{xy}(S_x^2 + S_y^2)\approx -S_z^4$,21,22 where $D_{CF}$ is the zero-field splitting parameter, or a Dzyaloshinsky-Moriya interaction (DMI) (Refs. 22 and 23) $D_{DMI}S_x\times S_y$. The DMI term has shown to be symmetry allowed on the kagome lattice and has been argued to prevail over the CF anisotropy in the spherical ($L = 0$) 3$d^5$ Fe$^{3+}$ ion.23 Fits to the spin-wave dispersion curves measured with neutron spectroscopy on single crystal24 and powder samples25 of potassium jarosite have found to be slightly better in a model with DMI compared to the model with CF anisotropy. The evi...
dence so far has, however, been inconclusive.

Given the similarity in crystal structure of potassium (iron) jarosite and hydronium (iron) jarosite the above effects could be expected to be equally important in the latter. However, hydronium jarosite has a spin-glass ground state with a freezing temperature of \( T_g = 17(2) \) K. The Weiss temperature for hydronium jarosite is less accurately known \((-700\to-1400 \) K) because the inverse susceptibility does not enter a linear regime sufficiently far below the decomposition temperature of the compound. It has been suggested that the spin-glass state is the result of proton transfer from the hydronium groups to the \( \text{Fe}-(\text{O}_6\text{H})\text{-Fe} \) superexchange mediating hydroxy groups. One indication that this could happen might be that the OH groups at the crystallographically analogous location in the related kagome compound zinc parataticamite can form a muonium state by chemically binding a positive muon. The aim of the work described here is to investigate the origins of the magnetic anisotropy (of either single-ion or DMI character) in the iron jarosites and to understand the essential differences between the salts, such as potassium jarosite, with a long-range-ordered ground state and hydronium jarosite.

Using Mössbauer spectroscopy the spin state of the \( \text{Fe}^{3+} \) in the iron jarosites was found to be \( S = 5/2 \), thus nominally high-spin \( 3d^5 \). The free-ion \( 3d^5 \) state is \( ^5S \), i.e., \( L = 0 \). Hence, there is no orbital angular momentum due to the spin-orbit coupling in the presence of a crystal field as long as the system is high-spin \( 3d^5 \). This holds at least up to third-order perturbation theory of the trigonal crystal-field distortions, spin-spin, and spin-orbit couplings. Even in a \( D_{4h} \) crystal field, the \( 3d^5 \) \( S = 5/2 \) spin should therefore be practically isotropic apart from the possible effects of dipole-dipole interactions. Both DMI and single-ion anisotropy arise only when the spin-orbit operator \( \mathbf{L} \cdot \mathbf{S} \) can induce a finite orbital angular momentum. That \( \langle L \rangle > 0 \) in the iron jarosites is clear from Curie-Weiss fits to the high-temperature magnetic susceptibility which indicate \( \mu_{\text{eff}} \) between 6.2 and 6.7 \( \mu_B \) in potassium iron jarosite and 6.6(2) \( \mu_B \) in the hydronium analog (compared to a spin-only value of the effective moment of 5.92 \( \mu_B \)). Clearly, this opens up the possibility for both DMI and single-ion anisotropies. However, the origin of the orbital angular momentum itself has so far received little attention.

II. EXPERIMENTAL

Single crystals of potassium iron jarosite \( \text{KF}_{2}(\text{OH})_{6}+(\text{SO}_{4})_{2} \) were grown using a hydrothermal reduction-oxidation method as described in Ref. 17: 4.88 g (28.0 mmol) \( \text{K}_2\text{SO}_4 \) and 2.2 mL (40 mmol) \( \text{H}_2\text{SO}_4 \) were dissolved in 50 mL distilled water and transferred into a 125 mL polytetrafluoroethylene (PTFE) liner of a stainless-steel bomb. 0.56 g (10 mmol) iron wire with a diameter of 2 mm was added to the solution. The bomb was placed in an oven at 202 °C for four days then cooled down to room temperature at a rate of 0.3 °C/min. The precipitate was washed, filtered, and dried, yielding 0.37 g, which is 22% based on Fe. In order to obtain sufficiently large crystals, it was important that the surface area of the solution exposed to air inside the vessel was minimized. This was achieved using a glass container which fitted inside the PTFE liner, with only a small hole at the top. Using this method single crystals of up to 0.7 mm in diameter were obtained, which was sufficiently large for the application in polarized x-ray spectroscopy. A number of crystals were characterized using single-crystal x-ray diffraction. These crystals were identified as potassium iron jarosite and face indexed.

The original solvothermal synthesis method for the jarosites was used for the preparation of \( \text{H}_3\text{OF}_{2}(\text{OH})_6+(\text{SO}_{4})_{2} \). In this preparation 6.6 g (22 mmol) of \( \text{Fe}_2(\text{SO}_4)_3 \cdot 5\text{H}_2\text{O} \) was dissolved in 50 mL water. This solution was transferred to a 125 mL PTFE liner of a stainless-steel bomb. The solution was heated in the bomb to 140 °C for 12 h. The hydronium iron jarosite which precipitated during the reaction was washed, filtered, and dried, yielding \( \sim 0.27 \) g of product.

Polarized soft x-ray absorption measurements were carried out using the liquid-helium cryostat housed in the high-field superconducting magnet on ID08 at the European Synchrotron Radiation Facility (ESRF) in Grenoble, France. Total-electron-yield spectra were measured by recording the drain current from the sample as a function of photon energy in the region of the Fe \( L_2,3 \) absorption edges. Potassium iron jarosite is strongly insulating, so the resonance enhanced total-electron-yield signal at the iron edge is much smaller.
FIG. 2. (Color online) X-ray absorption spectra of potassium iron jarosite at 290 K measured with x-ray polarization parallel and perpendicular to the c axis, $I_c$ (black circles), and $I_{ab}$ (red crosses), respectively. The lower trace shows the corresponding values of the isotropic absorption $I_{iso} = \frac{1}{2}I_{ab} + \frac{1}{2}I_c$ (black line) and the x-ray linear dichroism $I_{XLD} = I_{ab} - I_c$ (thin black line). Also shown $I_{iso}$ from a powder sample measured on 5U.1 at SRS Daresbury with a lower-energy resolution (broken red line). The inset shows a close up of $I_c$ and $I_{ab}$ at the $L_3$ edge.

than the signal from the surrounding metal of the sample holder. To avoid this source of background intensity, a potassium iron jarosite single crystal of 0.7 mm diameter was attached using silver glue to the end of a graphite tip, which in turn was glued to the end of a standard ESRF sample holder. The sample was aligned in the x-ray beam with the crystallographic c axis pointing in the horizontal plane and perpendicular to the incident beam. This allowed us to measure absorption spectra with the incident x-ray polarization both perpendicular and parallel to the c axis. Spectra were recorded with alternating horizontal and vertical polarizations and for each polarization averages were taken over at least eight scans. In this way the polarization-dependent spectra were taken at a number of temperatures between 20 and 290 K, in zero external field.

Isotropic absorption spectra at the Fe $L_{2,3}$ edges were obtained on beamline 5U.1 of the Synchrotron Radiation Source (SRS) at Daresbury Laboratory, U.K., using potassium and hydronium iron jarosite powder dispersed over double-sided UHV-compatible carbon tape.

III. RESULTS

Figure 2 shows the spectra obtained from single-crystal and powder samples of potassium iron jarosite at 290 K. The upper curves in Fig. 2 display the x-ray absorption spectra (XAS) from a single crystal measured on beamline ID08 at the ESRF, with the incident x-ray polarization parallel to the c axis $I_c$ (black circles) and perpendicular to the c axis, $I_{ab}$ (red crosses). The curves in the middle of the figure give the corresponding isotropic absorption spectrum $I_{iso} = \frac{1}{2}I_{ab} + \frac{1}{2}I_c$ (black line) and the isotropic spectrum directly measured from a powder sample on station 5U.1 at the SRS (broken red line). The curve at the bottom gives the x-ray linear dichroism $I_{XLD}$ defined as $I_{ab} - I_c$ (thin black line). The $2p_{1/2}$ transition consists of two edges; the first one at $\sim 709$ eV ($L_3$ edge) corresponds to the creation of a $2p_{3/2}$ core hole. The transition giving rise to a $2p_{1/2}$ core hole ($L_2$ edge) is at $\sim 12$ eV higher photon energy due to the $2p$ spin-orbit coupling. The inset of Fig. 2 shows the $I_c$ and $I_{ab}$ spectra in close up at the $L_3$ edge, which is the most significant region of the spectrum.

As seen in Fig. 3, the structure at the $L_3$ edge becomes more prominent in the isotropic spectrum as the temperature is lowered. At each stabilized temperature, the reproducibility of up to eight spectra demonstrated the absence of any temporal variation in the spectral line shape or intensity; therefore charging effects and drift in sample alignment can be eliminated as the cause of the observed temperature dependence. These isotropic spectra obtained by weighted averaging of $I_{ab}$ and $I_c$ are in good agreement with the spectra from powder samples measured for the SRS at all temperatures. That this change in line shape occurs in the powder averaged spectrum might be an indication of a slight change.
magnetic moments can be aligned by applying an external field, the x-ray magnetic circular dichroism (XMCD) and x-ray linear dichroism (XMLD) sum rules can be applied to obtain the expectation values of the orbital and spin magnetic moments respectively and the magnetocristalline anisotropy, respectively. In the case of the jarosites it is not possible to align the magnetic moments parallel with an external field due to the large energy scale of the antiferromagnetic interactions (T_CW = -800 K). The only way to obtain accurate information about the relevant expectation values is by explicitly calculating the transition probabilities,

\[ f(E_{h\alpha}, \mathbf{q}) = \frac{\langle \psi_{\alpha} | \mathbf{r}_{\mathbf{q}} | \psi_{\alpha} \rangle^2}{E_{h\alpha} - \Delta E - i\Gamma/2} \]  

between the ground state \( \psi_{\alpha} \) and excited state \( \psi_{\alpha'} \), where \( \mathbf{r}_{\mathbf{q}} \) is the electric-dipole operator for x-ray absorption with polarization \( \mathbf{q} \), and \( \Gamma \) is the lifetime broadening, which is treated as a fitting parameter. The experimental x-ray spectra were used to obtain good approximations for \( \psi_{\alpha} \) and \( \psi_{\alpha'} \) with the aid of the HILBERT+ code. The Hilbert space for \( \psi_{\alpha} \) is spanned by determinants of the lowest atomic LSJ configurations of \( |3d^5\rangle \) and \( |3d^{p+1}L\rangle \), where \( L \) denotes a ligand hole. Likewise, the Hilbert space for \( \psi_{\alpha'} \) is spanned by determinants of the atomic configurations of \( |2p^33d^{p+1}\rangle \) and \( |2p^33d^{p+2}L\rangle \). The Slater integrals for the 3d-3d and 2p-3d interactions and the 2p and 3d spin-orbit parameters for the ground state and final-state configurations calculated using Cowan’s atomic multiplet program are tabulated in Ref 52. Approximate solutions for the eigenfunctions of the iron ion in the compound are then obtained by diagonalization of the Hamiltonian including charge-transfer and crystal-field terms arising from the surrounding oxygen ligands,

\[ H = H_{\text{atom}} + \sum_b \left[ t_{\sigma,\pi,b} \langle \dot{z} \rangle_{\sigma,-} \langle p_{\pi} \rangle_{\sigma,+} + t_{\pi,\sigma,b} \langle d_{\sigma}^{3\pi} \rangle_{\sigma}\langle d_{\sigma}^{p} \rangle_{\sigma} + \text{c.c.} \right] + \sum_b \left[ V_{\sigma,\pi,b} \langle \dot{z} \rangle_{\sigma,-} \langle d_{\sigma}^{p} \rangle_{\sigma} + V_{\pi,\sigma,b} \langle d_{\sigma}^{3\pi} \rangle_{\sigma} \langle d_{\sigma}^{p} \rangle_{\sigma} + \text{c.c.} \right] + \frac{1}{2} \sum_b \langle p_{\pi}^{\sigma} \rangle_{\sigma} + \langle p_{\pi}^{\pi} \rangle_{\sigma} + \langle p_{\pi}^{\pi} \rangle_{\sigma} + \langle p_{\pi}^{\pi} \rangle_{\sigma}. \]

In this Hamiltonian, the first term represents the atomic Hamiltonian. The second term gives the hybridization between metal 3d and oxygen 2p orbitals, where \( t_{\sigma,\pi,b} \) and \( t_{\pi,\sigma,b} \) are the Slater-Koster hopping parameters. For each bond, \( t_{\sigma,\pi,b} \) is rescaled with the bond length using \( t_{\sigma,\pi,b} \rightarrow t_{\sigma,\pi,b} \left( R_{\text{ref}} / R_{\text{ref}} \right)^{\alpha} \) where \( R_{\text{ref}} \) is the normalized length of Fe-O bond \( b \) and the rescaling exponent \( \alpha \) is treated as a free parameter. The third term gives the contribution of the electrostatic crystal field which was not used in our case. The last term gives the energy of the 2p electrons in the oxygen valence band, where \( \epsilon_{p} \) is a simulation parameter related to the charge-transfer gap \( \Delta_{p} \) via \( \Delta_{p} = U_{3d,p} - \epsilon_{p} \). The total \( dd \) Coulomb energy for the addition of a sixth electron in the Fe 3d shell \( U_{3d,p} = \sum_{n=1}^{5} U_{n} \) was found to be 44.3 eV by considering the case \( \epsilon_{p}=0 \), where no charge transfer occurs unless the Fe\(^{2+}\) \( \Delta_{p} \) configuration has a lower energy than the Fe\(^{3+}\) \( \Delta_{p} \) configuration. The sum is made over all bonds \( b \), where the local coordinate frame \( \bar{x} \), \( \bar{y} \), and \( \bar{z} \) is oriented with the \( \bar{z} \) axis

\[ \int I_{\text{XMLD}}(\omega) d\omega \int I_{\text{XMLD}}(\omega) d\omega, \]  

FIG. 4. (Color online) Temperature dependence of the x-ray linear dichroism measured at the Fe \( L_{2,3} \) edges from a synthesized potassium iron jarosite single crystal (black lines) compared to calculated linear dichroism corresponding to the isotropic simulated spectra of Fig. 3 scaled in intensity to fit the experimental spectra by factors ranging from 1/2 at 290 K down to 1/3 for the 60 and 40 K data. Possible reasons for this mismatch are discussed in the text.

in the electronic structure as the temperature is lowered. Only very small changes were observed in the XLD spectral line shape on cooling the sample, as shown in Fig. 4. This change in the spectral line shape is accompanied by a slight reduction in the integrated XLD spectrum, 

\[ \int I_{\text{XMLD}}(\omega) d\omega \int I_{\text{XMLD}}(\omega) d\omega, \]
along the bond direction. Numerical comparisons between experiment and theory\textsuperscript{53,55} have shown that this approach works well provided that the Slater integrals for the Coulomb interactions and spin-orbit couplings are reduced to 70%–80% of the atomic Hartree-Fock values\textsuperscript{34} Here, all pd and dd Slater integrals were reduced to 80% of the atomic values calculated using Cowan’s program. The lifetime broadening $\Gamma$ was set to 0.24 eV for the $L_3$ edge and 0.3 eV for the $L_2$ edge. These are values as commonly used in the calculation of x-ray absorption spectra, with $L_2$ always slightly larger than $L_3$. This spectral broadening is due to the coupling with the (infinite dimensional) electromagnetic field.

The crystal field at the iron atom in potassium iron jarosite is approximately of $D_{4h}$ symmetry, with an elongation of the bond length $R_{ap}$ along the fourfold symmetry axis ($c'$) of $\sim 4\%$ with respect to the bond length $R_{eq}$ in the equatorial plane.\textsuperscript{17} The local symmetry axes of the FeO$_6$ octahedra are canted with respect to the crystallographic $c$ axis by an angle of $19.2\degree$, resulting in buckled FeO$_4$ planes forming the kagome layers.

Our simulation shows that the spectra are mainly determined by the strongly anisotropic $\sigma$-type hybridization between the iron 3d and oxygen 2p orbitals. The electrostatic part of the crystal field could be left out. An excellent agreement between experimental and simulated spectra for $I_{iso}$ was obtained using only four simulation parameters, namely, the charge-transfer gap $\Delta_{pd}$, $t_\sigma$, $t_\pi$, and the rescaling constant $\alpha$ as defined in $t_\sigma^2=\frac{R_{eq}}{R_{ap}}t_\sigma^2$ (with $R_{eq}/R_{ap}=1.035$ in the present case). When the simulated isotropic spectra are in good agreement with the experimental spectrum, at the same time optimal agreement is obtained for the linear dichroism spectrum. At high temperatures there is good agreement between the experimental and simulated linear dichroism spectrum, apart from an intensity scaling factor of 0.5, as shown in Figs. 4 and 5. With the fitting parameters for the simulation there is no way to reduce the overall intensity of the linear dichroism without radically modifying the isotropic spectrum. For up to 6\% the intensity difference might be attributable to the local canting of the FeO$_6$ octahedra with respect to the $ab$ plane. In addition to this there is an inevitable loss of intensity in the XLD due to misalignments and sample imperfection. Reduced energy resolution can also lead to a reduction in the measured XLD. We conclude that the difference in overall intensity of the experimental and calculated $I_{XLD}$ is not due to fundamental shortcomings of the simulation. For reasons we do not quite understand, the agreement between theory and experiment as evident in the simulation. For reasons we do not quite understand, the agreement between experimental and simulated spectra at low temperatures, the overall agreement between experiment and simulation as obtained here is remarkably good. The parameters as found in our simulation are given in Table I (left-hand side). The large error bar on the value for the charge-transfer gap of 3.5(2.0) eV reflects the adjustment of the total $dd$ Coulomb energy on the iron site $U_{pd}^d=U_{pd}(\langle n \rangle - 5)$ to any increase in $\varepsilon_p$ by the transfer of additional electron weight into the Fe ion. This is reflected in the total error in $\langle n \rangle$ of $\sim 0.1$ electron.

The values found here for $\Delta_{pd}$ and $t_\sigma$ are similar to those found previously for other Fe$^{3+}$ compounds.\textsuperscript{43,54} Fe$_3$O$_5$, which is in many respects probably the compound most closely related to iron jarosite, was previously found to have a band gap of predominantly charge-transfer character of 2–2.5 eV.\textsuperscript{55} With an unusually small value for $t_\sigma$ of 0.3(2) eV, the lowest energy peak at 707.8 eV is reproduced in the simulated spectrum. The main peak at 709 eV splits into two peaks with the introduction of a trigonal distortion of the FeO$_6$ octahedron. A common value for the rescaling exponent $\alpha$ is 3 and the small value found here ($\sim 7.5$) is an indication that the charge anisotropy is larger than the crystal-field anisotropy as defined by the oxygen positions around the iron. The lowest-energy low-spin configuration, corresponding to the $^4F_{11/2}$ free-ion configuration, lies 1.2 eV higher in energy and is not found to mix into the ground state.

The single-ion anisotropy $D_s$ is obtained as the difference between the energy of the $S_z=\pm 1/2$ and $\pm 5/2$ levels and was found to be of the easy-plane type. This easy-plane anisotropy was also confirmed by rotating the spin-quantization axis to various directions with respect to the crystal field. In this way we have also attempted to obtain estimates for $E_{xy}$. The $D_{ab}$ symmetry of the ligand field was lowered to $C_2$ to model the actual positions of the oxygen anions around the cation, as obtained from crystallographical data.\textsuperscript{11,17} However, this had no effect on the agreement of the calculated
TABLE I. Results from simulations of the Fe $L_{2,3}$ spectra of potassium iron jarosite, with ligand-field multiplet calculations using the HILBERT++ code (Ref. 50). The left-hand side lists all the parameters varied in the simulation. The right-hand side gives the resulting expectation values for the calculated ground-state properties of the Fe ion. The errors indicate the range over which the parameters could be varied while maintaining a good agreement with the data. The right-hand side of the lowest panel gives the effective moment in the case of full orbital polarization, while the calculation indicates only a very weak orbital polarization of the added electron weight due to charge transfer. We use the convention that upper case $J$,$L$,$S$ represent operators and lower case $j$,$l$,$s$ represent the quantum numbers given by $(L^2)=(l(l+1))$, etc.

<table>
<thead>
<tr>
<th>Parameters</th>
<th>Expectation values</th>
</tr>
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<tbody>
<tr>
<td>$\Delta_{pd}$ (eV)</td>
<td>3.5(2.0)</td>
</tr>
<tr>
<td>$t_\pi$ (eV)</td>
<td>0.3(2)</td>
</tr>
<tr>
<td>$\varepsilon$ (eV)</td>
<td>3.0(2)</td>
</tr>
<tr>
<td>$\alpha$</td>
<td>$\sqrt{2}\approx 2.32$</td>
</tr>
<tr>
<td>From $t_{\sigma,\pi}$ and $\alpha$ (in eV)</td>
<td>Using $(L)$ we obtain</td>
</tr>
<tr>
<td>$t_{\sigma}^{0}$</td>
<td>3.0(2)</td>
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<tr>
<td>$t_{\sigma}^{2}$</td>
<td>2.3(2)</td>
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<tr>
<td>$t_{\pi}^{0}$</td>
<td>$&lt;0.5$</td>
</tr>
<tr>
<td>$t_{\pi}^{2}$</td>
<td>$&lt;0.4$</td>
</tr>
<tr>
<td>Using $\sqrt{L^{2}}$ $\approx 2.77(6)$</td>
<td>2.77(6)</td>
</tr>
</tbody>
</table>

From Table I, it is clear that the ground-state expectation values are significantly larger than zero. It is not expected that in this 3$d$ transition-metal compound $l$ and $j(=s+l)$ are good quantum numbers, but there is sufficient evidence that the 3$d$ transition-metal compound is significantly larger than zero. It is not expected that in this 3$d$ transition-metal compound $l$ and $j(=s+l)$ are good quantum numbers, but there is sufficient evidence that the 3$d$ transition-metal compound is significantly larger than zero.

V. DISCUSSION

The expectation values of the ground-state obtained in our calculation are given in Table I (right-hand side). The corresponding free-ion configuration is $d^5$S$_{5/2}$ with an additional electron weight of $-0.4$ electron that is mainly in the $(l^2-y^2)$ orbital. The spin-orbit coupling is much smaller than the $t_{\sigma}$ hopping bandwidth and hence the fraction of the electron transferred to the ligands to the Fe cation is only very weakly orbitally polarized. The expectation values are much smaller than the orbital angular momentum as inferred from high-temperature magnetic-susceptibility data, which indicate that $\mu_{\text{eff}}=6.5\mu_B$. As shown in the lower panel of Table I (right-hand side) the experimental value is remarkably close to the case where the added electron weight due to charge transfer is fully orbitally polarized. It is not expected that in this 3$d$ transition-metal compound $l$ and $j(=s+l)$ are good quantum numbers, but there is sufficient evidence that the 3$d$ transition-metal compound is significantly larger than zero. It is not expected that in this 3$d$ transition-metal compound $l$ and $j(=s+l)$ are good quantum numbers, but there is sufficient evidence that the 3$d$ transition-metal compound is significantly larger than zero. It is not expected that in this 3$d$ transition-metal compound $l$ and $j(=s+l)$ are good quantum numbers, but there is sufficient evidence that the 3$d$ transition-metal compound is significantly larger than zero. It is not expected that in this 3$d$ transition-metal compound $l$ and $j(=s+l)$ are good quantum numbers, but there is sufficient evidence that the 3$d$ transition-metal compound is significantly larger than zero. It is not expected that in this 3$d$ transition-metal compound $l$ and $j(=s+l)$ are good quantum numbers, but there is sufficient evidence that the 3$d$ transition-metal compound is significantly larger than zero.
expectation values for $\langle S_y^2 \rangle$ at each corresponding experimental temperature, arising from the thermal population of the $m_s$ levels in the $S=5/2$ multiplet. The gradient dependence of $\langle S_y^2 \rangle$ (Fig. 6) shows a gradual alignment of the spins into the kagome planes at temperatures well above the transition temperature to a long-range-ordered state at 64 K. This is in rough agreement with the degree of coplanarity of the spins as measured using neutron diffraction on large single crystals.13

The simulated linear dichroism spectra at the lowest temperatures, corresponding to states with the spins aligned in the equatorial plane (the states $|s,m_s\rangle=|2.5,\pm 0.5\rangle$), do not agree well with the experimentally obtained spectra as shown in Fig. 4; although some features in the temperature dependence of the XLD are reproduced in the simulated spectra. Without success we have attempted to improve on this situation by calculating the spectra for the Fe spin aligned along the $x'$ axis in the equatorial plane (i.e., $|s,m_s\rangle=|2.5,\pm 2.5\rangle$) which resembles more closely the actual ground state in potassium jarosite. The remaining disagreement could be in part due to experimental error due to sample imperfections and in part due to intrinsic limitations of the simulation.

It might be naive to expect that the ligand field can be inferred from crystallographic data alone. A first indication for this is the large value of $\alpha$ obtained here, which amplifies the effect of the crystallographic trigonal distortion of the FeO$_6$ octahedra. Even when the symmetry of the ligand field is further lowered from $D_{4h}$ and accurately modeled on the locations of the oxygen anions around the central iron,37 neither the calculated spectra nor the simulation parameters change significantly. Therefore, it is possible that the actual charge anisotropy follows more closely the $xy$ plane rather than the equatorial oxygens. The easy-plane magnetic anisotropy of $D_{xy}=0.5(1)$ meV found here (see Table I) is then not directly related to the trigonal distortion of the crystal field but more to the two-dimensional (2D) character of the electronic hopping and magnetic exchange, the virtual absence of $\pi$ symmetry hybridization, and the Fe $3d$ spin-orbit coupling strength. In that case the easy plane could coincide more closely with the kagome plane. The strong 2D character can in part be attributed to the sulfate groups separating the kagome layers. The formal valence of the sulfur cation is 6+ with the (Ne) electronic configuration, but this large electrostatic charge is likely to attract electron weight from the neighboring oxygen anions, including the oxygens apical in the FeO$_6$ octahedra. The hopping parameters $t_{\sigma,a}$ from these oxygens to the iron are then further reduced with respect to the in-plane hopping. This translates into the large value for $\alpha$ found here.

The large difference between $t_\sigma$ and $t_{\sigma,a}$ obtained here might be explained by the triangular arrangement of the neighboring FeO$_6$ octahedra. This dramatically lowers the symmetry of the local crystal field around the oxygen ligands and reduces the $\pi$ symmetry overlap of the oxygen 2$p$ orbitals with the iron 3$d$ orbitals. Due to the low symmetry of the oxygen sites, the symmetry of the crystal field can be expected to be even lower than indicated by the crystallographic positions of the anions. It is in this sense fortunate that the situation can be modeled so well by the anisotropic $\pi$ and $\sigma$ hopping whose ratio is equal for all six anions around the central iron. It could well be that the last differences between the calculated and experimental spectra, such as the low-temperature x-ray linear dichroism, some of the difference in the overall intensity of the XLD, and the missing bump just above 707 eV, could be brought into agreement by allowing a different balance between $t_\sigma$ and $t_{\sigma,a}$ for the apical oxygen anions.

In neutron spin-wave studies22,24,25 the crystal-field anisotropy scenario has been compared in close detail with that of DMI. In the present context such a discussion would require calculations on clusters containing two to three iron centers. Such a calculation would certainly be of great interest but has not been attempted here, since the present approach already accounts so well for the experimental observations on potassium iron jarosite. The present result closes the last missing causal link from electronic structure to collective magnetic ground state.

VI. CONCLUSION

The isotropic and x-ray linear dichroism spectra taken at the Fe $L_{2,3}$ edges in potassium iron jarosite indicate that an orbital angular momentum and zero-field splitting arise due to an anisotropic, mainly $\sigma$ type, charge-transfer from the oxygen ligands. This closely resembles the scenario described in Ref. 38 to explain the zero-field splitting as observed in Mn$^{2+}$ salts.

The zero-field splitting parameter of the crystal-field anisotropy of 0.5 meV as found here is in excellent agreement with the value found from fits of a crystal-field anisotropy model to the spin-wave spectrum in the ground state of potassium iron jarosite.22,24,25 However, the most important result obtained here is that we have verified in detail the mechanism by which this magnetic anisotropy arises, closing...
the remaining missing link between the electronic structure in potassium iron jarosite and its magnetic ground state. We have even measured a change in the electronic structure as the magnetic ground state becomes coplanar, with the change in the isotropic line shape over the relevant temperature range. Of course, an additional in-plane anisotropy $E_{xy}$ and further-neighbor interactions are needed to fully explain the magnetic ground state. The present study does not provide more hints to the origin of the glassy state in hydronium jarosite. We found that the energy scale of the magnetic interaction in hydronium jarosite should be comparable to that in potassium jarosite. It might be that temperature-dependent spectra taken at a higher resolution could provide different insights in the case of hydronium jarosite. A final interesting observation is that the maximum orbital angular-momentum polarization possible for the amount of charge transfer observed here, but not the actually calculated orbital polarization, is in rough agreement with the experimentally observed effective moment.

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30 It has been shown (Ref. 32) that in fourth-, fifth-, and sixth-order perturbation theories of the crystal-field and spin-spin interactions a small orbital angular momentum and single-ion anisotropy arises.

31 T. Moriya, Phys. Rev. 120, 91 (1960).


Iron jarosite is in this sense not a unique case. For 3d transition-metal compounds with more than half-filled shells, the effective moments are in general significantly larger than the spin-only moment.

Though all Fe-O$_{eq}$ bond lengths are the same, the equatorial plane is rectangular rather than square, with the short side pointing into the kagome triangles. Furthermore, the Fe-O$_{ap}$ axis is canted by 4° away from the local $c'$ axis.