Absence of Spin Frustration in the Kagomé Layers of Cu²⁺ Ions in Volborthite Cu₃V₂O₇(OH)₂·2H₂O and Observation of the Suppression and Re-Entrance of Specific Heat Anomalies in Volborthite under an External Magnetic Field

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Abstract: We determined the spin exchanges between the Cu²⁺ ions in the kagomé layers of volborthite, Cu₃V₂O₇(OH)₂·2H₂O, by performing the energy-mapping analysis based on DFT+U calculations, to find that the kagomé layers of Cu²⁺ ions are hardly spin-frustrated, and the magnetic properties of volborthite below ~75 K should be described by very weakly interacting antiferromagnetic uniform chains made up of effective S = 1/2 pseudospin units. This conclusion was verified by synthesizing single crystals of not only Cu₃V₂O₇(OH)₂·2H₂O but also its deuterated analogue Cu₃V₂O₇(OD)₂·2D₂O and then by investigating their magnetic susceptibilities and specific heats. Each kagomé layer consists of intertwined two-leg spin ladders with rungs of linear spin trimers. With the latter as acting as S = 1/2 pseudospin units, each two-leg spin ladder behaves as a chain of S = 1/2 pseudospins. Adjacent two-leg spin ladders in each kagomé layer interact very weakly, so it is required that all nearest-neighbor spin exchange paths of every two-leg spin ladder remain antiferromagnetically coupled in all spin ladder arrangements of a kagomé layer. This constraint imposes three sets of entropy spectra with which each kagomé layer can exchange energy with the surrounding on lowering the temperature below ~1.5 K and on raising the external magnetic field B. We discovered that the specific heat anomalies of volborthite observed below ~1.5 K at B = 0 are suppressed by raising the magnetic field B to ~4.2 T, that a new specific heat anomaly occurs when B is increased above ~5.5 T, and that the imposed three sets of entropy spectra are responsible for the field-dependence of the specific heat anomalies.

Keywords: volborthite; deuterated volborthite; kagomé layer; spin exchange; DFT calculations; two-leg spin ladder; spin 1/2 antiferromagnetic uniform Heisenberg chain; entropy driven magnetic phase transitions; magnetic heat capacity; magnetic field-dependence of specific heat

1. Introduction

Properties of a magnetic material are described by a spin Hamiltonian, defined in terms of a few spin exchange paths between magnetic ions. The repeat pattern of strong spin exchange paths forms a spin lattice, with which the spin Hamiltonian generates the energy spectrum to be utilized in describing the magnetic properties. Consequently, the use of a correct spin lattice is paramount because the nature of the energy spectrum generated by a spin Hamiltonian depends on the nature of the spin lattice chosen. Antiferromagnets possessing a kagomé arrangement of transition-metal magnetic ions M are often believed to be spin-frustrated, hence suppressing a long-range magnetic order, so they are prime candidates that can give rise to exotic magnetic ground states [1,2]. However, spin frustration
is not guaranteed per se by, for example, a trigonal, a kagomé or a pyrochlore structural arrangement of magnetic transition-metal ions M. This is so because they interact strongly with the surrounding main-group ligands L to form MLₙ polyhedra, thereby splitting its d-states. What is crucial for geometrical spin frustration is not the geometrical pattern imposed by the arrangement of magnetic ions M, but rather that of the spin exchanges these ions generate with their neighboring ions. The direction-dependence of spin exchanges between magnetic ions M is determined by their magnetic orbitals, namely, the singly occupied d-states of their MLₙ polyhedra [3,4]. The d-states have the d-orbitals of M combined out of phase with the p-orbitals of the ligands L, so they are highly anisotropic in shape. Consequently, even if the magnetic ions form a kagomé arrangement, the spin exchanges between magnetic ions do not necessarily generate the same pattern. The anisotropy of spin exchange is most strongly manifested for a magnetic ion possessing only one magnetic orbital (e.g., S = 1/2 ions Cu²⁺ and Ti³⁺), and least strongly for a magnetic ion with five magnetic orbitals (e.g., S = 5/2 ions Mn²⁺ and Fe³⁺).

Volborthite, Cu₃V₂O₇(OH)₂·2H₂O, consisting of Cu²⁺ (d⁹, S = 1/2) ions in kagomé arrangement, has received much attention over the past decade [5–11], largely because it is believed to be spin-frustrated [5,11] The kagomé layers of Cu²⁺ ions (see Figure 1) are slightly distorted from an ideal kagomé shape with two different crystallographic sites for Cu²⁺ in a ratio 2:1. Below room temperature, each Cu²⁺ ion forms an axially elongated CuO₆ octahedron, so that, with the local z-axis taken along the elongated Cu-O bonds, the magnetic orbital of each CuO₆ octahedron is the x²−y² state contained in the CuO₄ equatorial plane, which is quite anisotropic in shape. In volborthite, the kagomé layers of composition Cu₃O₆(OH)₂ parallel to the ab-plane are pillared by pyrovanadates V₂O₇. The voids between the layers provide space for crystal water molecules. Below room temperature, volborthite undergoes two structural phase transitions, one at ~292 K from a C2/c phase to a I2/a phase, and the other at about ~155 K from the I2/a phase to a P2₁/c phase [6]. The latter structural phase transition generates two slightly different kagomé layers, which are slightly different in structure. Below 1.5 K, volborthite exhibits magnetic order, indicated by two anomalies in the magnetic specific heat [10].

![Figure 1](image_url)

**Figure 1.** (a) The kagomé arrangement of Cu²⁺ ions (red and blue circles) in volborthite, where the labels 1–5 refer to the spin exchanges J₁–J₅, respectively (see Table 1). In each J₂ path, two Cu²⁺ ions make a Cu-O . . . V⁵⁺ . . . O-Cu bridge, while the remaining exchange paths do not. (b) Geometrical entanglement of adjacent two-leg spin ladders in a kagomé layer. Adjacent spin ladders are presented with different colors for ease of distinction. (c) A two-leg spin ladder with rungs of linear trimers defined by J₂ and legs defined by J₄, with all exchange paths antiferromagnetically coupled, where the red and blue circles represent the up-spin and down-spin Cu²⁺ ion sites, respectively. (d) A spin ladder arrangement in which (FF), (AF), (AA) and (FA)-coupled spin ladders occur consecutively.
On the basis of their DFT+U calculations, Janson et al. [11] described the magnetic properties of volborthite using a trigonal spin lattice made up of pseudo S = 1/2 units, i.e., linear spin trimers in which adjacent Cu$^{2+}$ ions are strongly coupled antiferromagnetically. However, with this spin lattice model, it is difficult to explain why volborthite undergoes a magnetic ordering below 1.5 K [10]. Furthermore, the magnetic susceptibility Janson et al. calculated for volborthite by their exact diagonalization method for the spin Hamiltonian of the trigonal lattice show a spin gap below the susceptibility maximum at ~17 K. The latter is in sharp contrast to the experimental observation [5], which reveals that the magnetic susceptibility remains rather high as the temperature is decreased toward 0 K [5]. Thus, results of Janson et al.’s exact diagonalization based on their trigonal spin lattice model are inappropriate for understanding the physics of volborthite below ~17 K.

The magnetic entropy removed by the long-range order is small compared to $R \ln 2$, as is typically found for low-dimensional short-range ordered (SRO) magnetic systems, e.g., for S = 1/2 antiferromagnetic uniform Heisenberg (AUH) chains. Then, one might speculate if the magnetic ordering of volborthite observed below 1.5 K is associated with a magnetic ordering of such S = 1/2 AUH chains, although it has been believed that volborthite is spin-frustrated [5,11].

To explore the magnetic order below 1.5 K, especially its dependence on the magnetic field, as well as the origin of the apparent finite susceptibility as T→0 K, we re-analyzed the spin lattice of volborthite by performing an energy-mapping analysis based on DFT+U calculations [3,4] to find that the magnetic properties of volborthite below ~75 K should be described by two-leg spin ladders with rungs of S = 1/2 pseudospin units rather than by a trigonal lattice of S = 1/2 pseudospin units, as proposed by Janson et al. [11]. We verified this conclusion by acquiring new magnetic susceptibility data and re-analyzing reported magnetization data of volborthite to show that the kagomé layer of Cu$^{2+}$ ions is hardly spin-frustrated, and the low-temperature magnetic properties of volborthite should be described by an AUH chain composed of S = 1/2 pseudospin units. We characterized the magnetic phase transitions below 1.5 K by carrying out specific heat measurements for Cu$_3$V$_2$O$_7$(OH)$_2$·2H$_2$O and its deuterated analogue, Cu$_3$V$_2$O$_7$(OD)$_2$·2D$_2$O, under a magnetic field B = 0–9 T. Our work shows that the magnetic ordering of volborthite below 1.5 K is suppressed by field B > ~4.5 T, while a new magnetic ordering takes place when B > ~5.5 T, and that these field-dependent behaviors of the magnetic ordering originate from the three sets of magnetic entropy spectra of each kagomé layer of Cu$^{2+}$ ions, created by constrained interactions between adjacent two-leg spin ladders.

2. Experimental and Calculations

To determine the spin exchanges of the I2/a and P21/c phases of volborthite, we carried out DFT+U calculations, employing the Vienna ab Initio Simulation Package (VASP) [12,13] using the projector augmented wave (PAW) [14,15] method and the PBE [16] exchange-correlation functional. The electron correlation associated with the 3d states of Cu was taken into consideration by DFT+U calculations, i.e., by performing DFT calculation with an effective on-site repulsion $U_{\text{eff}} = U - J = 4$ and 5 eV added on the magnetic ions Cu$^{2+}$ [17].

Single crystals of volborthite were grown using hydrothermal techniques, as described in the literature [6]. Deuterated samples of volborthite were prepared by replacing light by heavy water (isotope enrichment 99.5%). The magnetic susceptibilities were measured at constant field as a function of the temperature in a Magnetic Property Measurement System SQUID magnetometer (MPMS-XL7, Quantum Design, San Diego, CA, USA). The specific heats of a collection of oriented crystals were determined using the relaxation method of a 3-He Physical Property Measurement System (PPMS, Quantum Design, San Diego, CA, USA). In order to construct a lattice reference needed to evaluate the magnetic contribution to the specific heat of volborthite, we prepared a polycrystalline sample of the diamagnetic mineral martyrte with composition Zn$_3$V$_2$O$_7$(OH)$_2$·2H$_2$O from an aqueous solution of NH$_4$VO$_3$ and zinc acetate, Zn(CH$_3$CO$_2$)$_2$, following a recipe reported in the literature [18].
3. Results

3.1. Spin Exchanges and Spin Lattice of Volborthite

We determine the spin exchanges $J_1$-$J_5$ defined in Figure 1a by performing the energy-mapping analyses [3,4] based on DFT+U calculations (for details of calculations, see the Supplemental Material). Below room temperature down to ~155 K, volborthite has a crystal structure with two equivalent kagomé layers in the unit cell (space group $I2/a$) and below 155 K a structure with two slightly non-equivalent kagomé layers of Cu$^{2+}$ ions (space group $P2_1/a$) [5,8]. The spin exchanges of the two phases are very similar, as summarized in Table 1, where only the values obtained with $U_{\text{eff}} = 4$ eV are listed. Those calculated with $U_{\text{eff}} = 5$ eV are given in the Supplemental Material. Janson et al. [11] carried out DFT+U calculations using $U_{\text{eff}} = 8.5, 9.5$ and 10.5 eV to determine the spin exchanges of the $P2_1/a$ structure by fitting the electronic band structure in terms of a tight binding approximation.

In the following, we first discuss the spin lattice of volborthite suggested by the spin exchanges we obtained. Subsequently, we show that the same picture is obtained from those of Janson et al., despite that their spin exchanges are considerably smaller than ours in the strengths of AFM character due to their use of very large $U_{\text{eff}}$ values.

As can be seen from Table 1, the strongest exchange $J_2$ is AFM and forms isolated linear trimers (Figure 1a). The second strongest spin exchange $J_4$ is also AFM but is weaker than $J_2$ by nearly an order of magnitude. All other spin exchanges are weaker than $J_2$ by more than an order of magnitude. Since $J_2$ is much stronger than other spin exchanges, each linear trimer at $T << J_2$ constitutes a pseudospin $S = 1/2$ unit and such units form a triangular lattice, as already pointed out by Janson et al. [11] However, we note that the linear trimers interact through the AFM exchange $J_4$ to form two-leg spin ladders with the linear trimers as rungs (Figure 1b). This feature is hidden in a kagomé layer because nearest-neighbor spin ladders are geometrically entangled with their superposed legs.

The two-leg spin ladder model as the spin lattice of volborthite described above is also supported by Janson et al.’s spin exchanges (Table 1), although their spin exchanges differ considerably from ours in magnitude. In general, a spin exchange $J$ between two magnetic ions located at sites $i$ and $j$ can be written as the sum of the FM and AFM components ($J_F$ and $J_{AF}$, respectively), namely, $J = J_F + J_{AF}$ [3,4,19]. With the magnetic orbitals describing the spin sites $i$ and $j$ as $\psi_i$ and $\psi_j$, respectively, the overlap density $\rho_{ij} = \psi_i^\dagger \psi_j$ gives rise to the exchange repulsion $K_{ij}$, while the overlap integral $<\psi_i^\dagger \psi_j>$ leads to an energy split $\Delta e$ between the two states described by the magnetic orbitals. Then, $J_F$ and $J_{AF}$ are written as

$$J_F = -K_{ij}, \quad \text{and} \quad J_{AF} = \frac{(\Delta e)^2}{U_{\text{eff}}} \quad (1)$$

Since the AFM component $J_{AF}$ is inversely proportional to $U_{\text{eff}}$, using a large $U_{\text{eff}}$ value in DFT+U calculations should make $J$ value less AFM or even shift it to a FM spin exchange. This explains why the $J_2$ and $J_4$ values of Janson et al. are less strongly AFM than ours, and why their $J_1$ and $J_3$ values are more strongly FM than ours.

Table 1. Values (in K) of the spin exchanges $J_1$–$J_5$ calculated for the $I2/a$ and $P2_1/a$ phases of volborthite obtained by DFT+U calculations.

| $I2/a$ Phase $^a$ | $P2_1/a$ Phase $^a$ | $P2_1/a$ Phase $^b$ |
|-------------------|---------------------|---------------------|
|                   | Layer 1 | Layer 2 | Layer 1 | Layer 2 | Layer 1 | Layer 2 |
| $J_1/J_2$         | 0.010   | −0.001  | −0.025  | −0.15   | −0.11   |
| $J_2/J_2$         | 0.025   | 0.031   | 0.031   | −0.34   | −0.32   |
| $J_4/J_2$         | 0.145   | 0.141   | 0.135   | 0.17    | 0.15    |
| $J_5/J_2$         | 0.014   | 0.014   | 0.013   | -       | -       |
| $J_2$             | 542 K   | 550 K   | 582 K   | 193 K   | 205 K   |

$^a$ Present work with $U_{\text{eff}} = 4$ eV. $^b$ Janson et al. (ref. [11]) with $U_{\text{eff}} = 8.5$ eV.
We now examine the interactions between adjacent two-leg spin ladders, under the constraint that the spin exchanges \( J_2 \) and \( J_4 \) forming the two-leg spin ladders are much stronger than \( J_3 \) and \( J_1 \), which provide interactions between spin ladders. The latter requires that, in all possible spin ladder arrangements, all \( J_2 \) and \( J_4 \) exchange paths of each two-leg spin ladder must remain antiferromagnetically coupled (Figure 1d). In every intertwined leg, each \( J_2 \) leg has two consecutive \( J_3 \) paths. Since each \( J_2 \) leg is antiferromagnetically coupled, \( J_3 \) has no effect on the interaction between adjacent two-leg spin ladders, regardless of whether it is AFM or FM. Thus, adjacent two-leg spin ladders interact only through the very weak exchanges \( J_1 \) (Figure 1a). Note that \( J_1 \) is weakly AFM for the \( I2/a \) phase but is weakly FM for the \( P2_1/a \) phase in our calculations. This difference does not influence how the ordering between spin ladders in a kagomé layer is affected by \( J_1 \) (see below). Since \( J_1 \) is more than an order of magnitude weaker than \( J_4 \), the magnetic property of each kagomé layer at low temperatures (\( |J_1| < T \ll J_2 \)) is primarily determined by those of the two-leg spin ladders with each ladder acting as an effective \( S = 1/2 \) pseudospin rung (Figure 1c). At sufficiently low temperatures where excitations within the rungs are negligible, the dominant AFM spin exchanges form two-leg spin ladders predicting that volborthite should be regarded primarily as a system of very weakly coupled \( S = 1/2 \) AUH chains. In the following, we first demonstrate that the magnetic properties of volborthite below \( \sim 75 \) K are well explained by the model of very weakly interacting \( S = 1/2 \) AUH chains.

3.2. Experimental Evidence for the Spin-Half AFM Uniform Heisenberg Chain Character

Figure 2 displays the magnetic susceptibilities of volborthite, which exhibits a broad peak at \( \sim 17 \) K and a finite susceptibility as \( T \to 0 \) K, characteristic of low-dimensional AFM SRO behavior. As implied by the results of the DFT+U calculations, the magnetic susceptibilities of volborthite between 3 and 75 K can indeed be very well fitted by those theoretically predicted for a \( S = 1/2 \) AUH chain according to

\[
\chi_{\text{mol}}(T) = \chi_{\text{spin}}(J_C, g, T) + \chi_0
\]

where \( \chi_{\text{spin}}(J_C, g, T) \) represents the magnetic susceptibility of the \( S = 1/2 \) AUH chain with nearest-neighbor (NN) spin exchange \( J_C \) (Figure 2a). For \( \chi_{\text{spin}}(J_C, g, T) \), we used the Padé approximant of Quantum Monte Carlo (QMC) results for the \( S = 1/2 \) AUH chain [20]. The second term refers to the temperature-independent susceptibility \( \chi_0 \) arising from the diamagnetism of the electrons in the closed shells \((-175 \times 10^{-6} \) cm\(^3\)/mol per formula unit, FU) [21,22] and also from the van Vleck paramagnetic susceptibility due to excitations to the excited states of each \( \text{Cu}^{2+} \) ion \((-100 \times 10^{-6} \) cm\(^3\)/mol per Cu atom) [23], leading to \( \chi_0 = +125 \times 10^{-6} \) cm\(^3\)/mol per FU. Up to \( \sim 75 \) K, the experimental susceptibility is well reproduced by the susceptibility calculated for \( J_C = 27.8(5) \) K and \( g = 2.33 \) (the solid blue curve in Figure 2a). The fitted \( g \)-factor is found to be 2.33, which is at the higher end of the \( g \)-factor expected for \( \text{Cu}^{2+} \) ions [24]. The \( S = 1/2 \) AUH chain model readily explains the susceptibility maximum at \( T_{\text{max}} = 17 \) K as well as the finite susceptibility below \( T_{\text{max}} \). The exchange \( J_C = 27.8(5) \) K agrees very well with the expected value \( J_C = T_{\text{max}}/0.64085 = 26 \) K. In addition, the ratio \( \chi_{\text{spin}}T_{\text{max}}/g^2 = 0.0346 \) cm\(^3\)/K/mol concurs well with the value 0.0353 expected for the \( S = 1/2 \) AUH chain [20]. The difference between the experimental data and the chain susceptibility (green solid line in Figure 1) vanishes below \( \sim 75 \) K and gradually increases to higher temperatures. The susceptibilities above \( \sim 75 \) K without any constraints matches very well with the susceptibility of an isolated linear spin \( S = 1/2 \) trimer described by the spin Hamiltonian

\[
H = J_{\text{trim}}(\vec{S}_1 \cdot \vec{S}_2 + \vec{S}_2 \cdot \vec{S}_3)
\]

where \( \vec{S}_i \) (\( i = 1, 2, 3 \)) represent the three spin sites. Replacing \( \chi_{\text{spin}}(J_C, g, T) \) in Equation (2) with the susceptibility of a spin \( S = 1/2 \) trimer given e.g., by Boukhari et al. [25], the susceptibility data above \( \sim 75 \) K can be well fitted without further constraints. The trimer
spin exchange amounts to 197(2) K, indicating a ratio $J_C/J_{\text{trim}} = 0.14$, in good agreement with the ratio $J_4/J_2$ obtained from our DFT+U calculation (see Table 1), suggesting that the spin trimers should be identified as the rungs of the two-leg spin ladders running along the crystallographic $b$ axis.

$$\text{Figure 2. (a) The magnetic susceptibility of one formula unit (i.e., comprising three Cu atoms) of volborthite (black circles) with } B \text{ applied along the crystallographic } b \text{ axis fitted (for } T < 75 \text{ K) to the theoretical prediction for a } S = 1/2 \text{ AUH chain (solid blue curve) given by Equation (2). The difference between the two is displayed as a solid green line. The experimental susceptibilities for } 75 \text{ K} \leq T \leq 320 \text{ K are well fitted by the susceptibility of a linear spin } S = 1/2 \text{ trimer with spin exchange of } 197(2) \text{ K (red dotted curve). Note the susceptibility anomaly at ~295 K due to the structural phase transition from the } C2/c \text{ to the } I2/a \text{ phase [3]. (b) Isothermal magnetization of volborthite determined at 1.4 K taken from ref. [11] compared with quantum Monte Carlo calculations for a } S = 1/2 \text{ AUH chain with a spin exchange of } J_C = 27.5 \text{ K (solid red line).}$$

Another characteristic feature of the magnetic properties of volborthite is the low-temperature 1/3 magnetization plateau extending over a wide magnetic field range, displayed in Figure 2b [8,9]. Note that we multiplied the published literature data by a factor of three to obtain the magnetization per one FU of volborthite (3 Cu atoms); the experimental data are well reproduced by the magnetization predicted for the $S = 1/2$ AUH chain with $J_C = 27.5$ K. The theoretical magnetization was obtained by Quantum Monte Carlo Calculations (QMC), employing the path integral method of the loop code incorporated within the ALPS project [26,27]. To ensure a low statistical error, these simulations included 250 spins and $1.5 \times 10^5$ Monte Carlo steps for the initial, as well as the subsequent, thermalization.

### 3.3. Specific Heat Anomalies of Volborthite
#### 3.3.1. Specific Heat Anomalies of Volborthite

In Figure 3a, we show the specific heat measured for an ensemble of aligned crystal of volborthite $\text{Cu}_3\text{V}_2\text{O}_7(\text{OH})_2\cdot2\text{H}_2\text{O}$ in zero magnetic field with magnetic field applied along the $b$ axis, the estimated lattice contribution to the heat capacity constructed from the specific heat of $\text{Zn}_3\text{V}_2\text{O}_7(\text{OH})_2\cdot2\text{H}_2\text{O}$ [28] and the difference between the two representing the magnetic contribution to the specific heat of volborthite. It is difficult to obtain a proper lattice specific heat reference because the atomic masses of Cu and Zn are not the same, and because the space groups describing the crystal structures of $\text{Zn}_3\text{V}_2\text{O}_7(\text{OH})_2\cdot2\text{H}_2\text{O}$ ($P\bar{3}m1$) and $\text{Cu}_3\text{V}_2\text{O}_7(\text{OH})_2\cdot2\text{H}_2\text{O}$ ($I2/a$ and $P2_1/a$) are different. In our work, we adopt the procedure suggested by Boo and Stout [29] and stretch the temperature axis of the specific heat data for $\text{Zn}_3\text{V}_2\text{O}_7(\text{OH})_2\cdot2\text{H}_2\text{O}$ with a smooth function, varying from 1.15 at 0 K to 1.05 at 50 K. The latter values were chosen such that the specific heat of $\text{Zn}_3\text{V}_2\text{O}_7(\text{OH})_2\cdot2\text{H}_2\text{O}$ smoothly approaches that of volborthite as $T \rightarrow 50$ K. The magnetic specific heat of volborthite thus obtained exhibits a broad maximum consistent with the specific heat capacity expected for a $S = 1/2$ AUH, with a NN spin exchange of $J_C = 26$ K. The latter value is in good agreement with the spin exchanges derived from the analyses of
the magnetic susceptibility and magnetization data. The total magnetic entropy amounts to ~80% of Rln2 expected for a S = 1/2 system, consistent with the result reported by Yamashita et al. [10].

![Figure 3](image-url) **Figure 3.** (a) Specific heats of volborthite (black dots) and martyite, Zn$_3$V$_2$O$_7$(OH)$_2$·2H$_2$O (red dots). The latter was scaled, as described in detail in the text, to smoothly approach the heat capacity of volborthite at high temperatures. The difference (blue dots) representing the magnetic contribution to the specific heat of volborthite. The solid red line displays the heat capacity of a S = 1/2 AUH with spin exchange of 26 K and spin entropy amounting to ~80% of Rln2 expected for a S = 1/2 system. (b) Comparison of the specific heats of Cu$_3$V$_2$O$_7$(OH)$_2$·2H$_2$O and Cu$_3$V$_2$O$_7$(OD)$_2$·2D$_2$O.

### 3.3.2. Effect of Isotope Substitution

In Figure 3b, we compare the specific heats of Cu$_3$V$_2$O$_7$(OH)$_2$·2H$_2$O and its deuterated analogue, Cu$_3$V$_2$O$_7$(OD)$_2$·2D$_2$O, prepared by using 99.5% isotope enriched heavy water in the preparation. The specific heat anomalies of Cu$_3$V$_2$O$_7$(OH)$_2$·2H$_2$O below 1.5 K (see inset Figure 3b) are hardly affected by the deuteration. However, the structural phase transition from C2/c to I2/a near 155 K undergoes a 5% upshift for the deuterated sample. This strongly suggests that the two specific heat anomalies below 1.2 K are related to magnetic ordering. Apparently, the D→H substitution does not essentially modify the spin exchange between adjacent kagomé layers, but substantially affects the structural properties of volborthite. The latter finding reflects the fact that the pyrovanadate groups connecting neighboring kagomé layers provide large interlayer spacing. The hydrogen bond network, from an OH group of a kagomé layer to a crystal water to an O atom of the adjacent kagomé layer, is loosened by the D→H substitution, which facilitates the I2/a to P2$_1$/a structural transition in Cu$_3$V$_2$O$_7$(OD)$_2$·2D$_2$O.

### 3.3.3. Effect of a Magnetic Field on Magnetic Ordering

Given that the specific heat anomalies below 1.5 K are associated with magnetic ordering, it is reasonable to expect that it can be destroyed, and the specific heat anomalies disappear under the sufficiently high external magnetic field. To confirm this conjecture, we carried out specific heat measurements for volborthite under magnetic fields B = 0–9 T (Figure 4a). As B increases, the two specific heat anomalies observed at B = 0 widen but remain separated, forming two ridges until they eventually merge into one and then abruptly disappear for B > ~4.2 T (see Figure 4a). Surprisingly, however, a new specific heat anomaly occurs for fields B > ~5.5 T. This new anomaly forms a single broad ridge, widening and shifting to higher temperatures with increasing B.
remain separated, forming two ridges until they eventually merge into one and then an anomaly occurs for fields $T$ (Figure 4a). As $K$, the free energy it can exchange energy with the surrounding phonon bath. At a temperature necessary to find the magnetic energy spectra of volborthite available below 1.5 K, with which we carried out specific heat measurements for volborthite under magnetic fields.

To probe the cause for the field-dependence of the specific heat anomalies, it is necessary to find the magnetic energy spectra of volborthite available below 1.5 K, with which it can exchange energy with the surrounding phonon bath. At a temperature $T$ below 1.5 K, the free energy $G_i$ of volborthite associated with its magnetic arrangement $i$ can be written as $G_i = H_i - TS_i$, where $H_i$ and $S_i$ are the enthalpy and entropy of the magnetic arrangement $i$, respectively. If volborthite has a set of different magnetic arrangements $i = 1, 2, 3, \ldots$, it has the corresponding energy spectrum $\{G_1, G_2, G_3, \ldots\}$. Then, volborthite can absorb energy from the surroundings by using this energy spectrum reflected by the specific heat anomaly. Since the magnetic specific heat anomalies are observed below 1.5 K, one might assume that the enthalpies $H_i$ are identical for all members $i = 1, 2, 3, \ldots$. Then, the free energy spectrum $\{G_1, G_2, G_3, \ldots\}$ is determined by the corresponding entropy spectrum $\{S_1, S_2, S_3, \ldots\}$, so the specific heat anomalies below 1.5 K arise from the entropy spectrum of volborthite, which is due most likely to the magnetic ordering of each kagomé layer. The latter implies that each kagomé layer must possess at least two different sets of entropy spectra, so that one set is responsible for the specific heat anomalies below ~4.2 K, and the other for those above ~5.5 K. In the next section, we show that each kagomé layer has three different sets of entropy spectra associated with the arrangements of two-leg spin ladders in each kagomé layer. These entropy spectra are a direct consequence of the fact that interactions between adjacent two-leg spin ladders in each kagomé layer are topologically constrained (see below).

4. Entropy Spectra and Specific Heat Anomalies

4.1. Spin Ladder Arrangements

In this section we discuss the magnetic-field dependence of the magnetic ordering in volborthite below 1.5 K. Our experiments gave conclusive evidence for an effective AUH chain behavior of volborthite at low temperatures, based on the two-leg spin ladders formed by $J_2$ and $J_4$. As already mentioned, the two-leg spin ladders (thereafter, referred to as the spin ladder for short) interact with their neighboring spin ladders to establish two-dimensional correlations within each kagomé layer via the spin exchange $J_1$ (Section 3.1). To begin with, we examine what kinds of spin ladder arrangements are possible in a kagomé layer. As already mentioned, in all possible magnetic arrangements of each kagomé layer at temperatures below 1.5 K, all $J_2$ and $J_4$ exchange paths of every spin ladder should remain antiferromagnetically coupled (Figure 1d) because the spin exchanges $J_2$ and $J_4$ forming the spin ladders are AFM and are much stronger in magnitude than $J_1$. This constraint on the magnetic arrangements of a kagomé layer leads to three different groups of spin ladder arrangements. Note that each spin ladder has a set of two consecutive $J_1$ paths running...
along the leg direction, which consists of the left and right sets of $J_1$ paths. Then, the first topological constraint is that the left or right set of $J_1$ paths in each spin ladder be coupled either all antiferromagnetically or all ferromagnetically. For simplicity, we use the labels A (F) to represent the set of $J_1$ paths all antiferromagnetically (ferromagnetically) coupled in each spin ladder. The two sets of the $J_1$ paths in each spin ladder can have one of the four possible arrangements, (AA), (AF), (FA), and (FF), as illustrated in Figure 1d. Here, the (AF) arrangement, for example, means that the spin ladder indicated by a parenthesis has the $J_1$ paths all antiferromagnetically coupled in the left set, but all ferromagnetically coupled in the right set. Use of bold labels in the (AA) and (FF) arrangements stems from the ease of recognizing the spin ladder arrangements, as will be seen below. Figure 1d illustrates a kagomé layer in which the (FF), (AF), (AA) and (FA) spin ladders occur consecutively. It is convenient to denote this arrangement of spin ladders by (FF)(AF)(AA)(FA). As can be seen from this arrangement, the second topological constraint is that, when crossing from one spin ladder to its adjacent one, the two sets of the $J_1$ paths adjacent to the boundary between the two be opposite in the nature of the spin exchange coupling, e.g., “F)(A” or “A)(F”. As a consequence, the third topological constraint is that (AA) spin ladders cannot be nearest neighbors, nor can be (FF) spin ladders. This implies also that (AA) and (FF) ladders always have to alternate in any spin ladder arrangement. These three constraints imply that there occur only three different sets of spin ladder arrangements in each kagomé layer, in which the members of each set are distinguished by how many pairs of (AA) and (FF) spin ladders are present. For simplicity, we use $(AF)_m$ to represent the occurrence of $m$ consecutive (AF) spin ladders, and $(FA)_n$ that of $n$ consecutive (AF) spin ladders. Then, with the integers $m$, $n$, $o$, $p$, $q$ and $r$ counting the numbers of consecutive (AF) or (FA) arrangements, examples of the spin ladder arrangements in Groups I, II and III are written as follows:

Group I: $(AF)_m(FA)_o(FF)(AF)_p(FA)_q(FF)(AF)_r$
Group II: $(AF)_m(FA)_o(FF)(AF)_p(FA)_q(FF)(AF)_r(FA)(FA)_r$
Group III: $(FA)_m(FF)(AF)_n(AA)(FA)_o(FF)(AF)_p(FA)_q(FF)(AF)_r$

The example of Group I given above illustrates the case when two pairs of (AA) and (FF) spin ladders are separated by patches of consecutive (FA) and (AF) spin ladders. All possible spin ladder arrangements of Group I are obtained by changing the number of (AA) and (FF) pairs, $m = \{0, 1, 2, 3, \ldots \}$. With respect to any member of Group I, the corresponding member of Group II has one more (AA) spin ladder, while that of Group III has one more (FF) spin ladder.

4.2. Entropy Spectra

For a given $(AF)_m(FA)_n$ arrangement, it is important to note that the $(AF)_m$ patch represents a domain of (AF)-coupled spin ladders while the $(FA)_n$ patch represents a domain of (FA)-coupled spin ladders, and the $(AA)$ spin ladder is a boundary between the two domains. Similarly, in a given $(FA)_m(FF)(AF)_n$ arrangement, the $(FF)$ spin ladder represents a boundary between the $(FA)_m$ and $(AF)_n$ domains. Thus, each member of the Groups I, II and III arrangements is characterized by how many pairs of (AA) and (FF) boundaries it has, regardless of the widths (e.g., values of $m$, $n$, $\ldots$) of each individual domain. This point is important in connection with the enthalpy of any given spin ladder arrangement. All members of Group I have an identical enthalpy, $H_1 = J_1 \sum_{i>j} \vec{S}_i \cdot \vec{S}_j = 0$ per kagomé layer, because each member has equal numbers of antiferromagnetically and ferromagnetically coupled $J_1$ paths. Similarly, all spin ladder arrangements of Group II are identical in enthalpy, $H_{II} = H_1 - 2J_1 = -2J_1$ per kagomé layer, and those of Group III are identical in enthalpy, $H_{III} = H_1 + 2J_1 = +2J_1$ per kagomé layer. Note that $J_1$ is weakly FM in the $P2_1/\alpha$ phase (see Table 1), and that the value of $-2J_1$ for Group II is the enthalpy of one (AA) spin ladder while the value of $+2J_1$ for Group III is the enthalpy of one (FF) spin ladder. Since $J_1$ is very weak and since a kagomé layer has a large number of spin ladders,
one can for all practical purposes assume \( H_I = H_{II} = H_{III} = 0 \) per kagomé layer, that is, all spin ladder arrangements of Groups I, II and III are practically identical in enthalpy.

The spin ladder arrangements of Groups I, II and III give rise to entropy spectra that are characterized by the number \( m \) of (AA) and (FF) pairs. To probe the entropy spectra of Groups I, II and III, we consider that a kagomé layer has \( N \) spin ladders, where \( N = N_A/3 \) per FU of volborthite \( \text{Cu}_3\text{V}_2\text{O}_7(\text{OH})_2\cdot2\text{H}_2\text{O} \) with \( N_A \) as Avogadro’s number. The factor 1/3 takes into account the fact that each spin ladder has three \( \text{Cu}^{2+} \) ions per repeat unit. For the case when there are \( m \) pairs of (AA) and (FF) arrangements, the total number of different spin ladder arrangements, \( \Omega(A)(2m) \), in Group I, is given by

\[
\Omega_I(2m) = 2(NC_{2m}) = 2(N!)/[(2m)!(N - 2m)!],
\]

where \( NC_{2m} \) is the binomial expansion coefficient, and the factor 2 arises because the \( m \) pairs of sites in each arrangement can be occupied first with a (FF) spin ladder or first with an (AA) spin ladder when \( m > 0 \). For \( m = 0 \), \( \Omega_I(0) = 2 \) because one can have the (AF)\(_\infty\) = \ldots (AF)(AF)(AF) \ldots or the (FA)\(_\infty\) = \ldots (FA)(FA)(FA) \ldots arrangement. The corresponding numbers for Group II (Group III) are given by

\[
\Omega_{II}(2m) = \Omega_{III}(2m) = \Omega_I(2m) \Omega_E(2m),
\]

where \( \Omega_E(2m) \) represents the number of possible sites available for an extra spin ladder after \( m \) pairs of (AA) and (FF) spin ladders are chosen from \( N \) spin ladders. Due to the topological constraints on the spin ladder ordering, \( \Omega_E(2m) \leq N - 2m \). Then, the entropies \( S \) associated with the three groups of spin ladder arrangements are given by

\[
S_I(2m) = k_B \ln \Omega_I(2m),
\]

\[
S_{II}(2m) = S_{III}(2m) = S_I(2m) + S_E(2m),
\]

\[
S_E(2m) = k_B \ln \Omega_E(2m).
\]

Since \( \Omega_I(2m) = \Omega_I(N - 2m) \), \( \Omega_I(2m) \) increases as 2m increases from 0 toward \( N/2 \), reaching a maximum at \( 2m = N/2 \), and decreases as 2m increases beyond \( N/2 \) toward \( N \) (Figure 4b). Consequently, \( S_I(2m) = S_I(N - 2m) \), and \( S_I(2m) \) increases as 2m increases from 0 toward \( N/2 \) and as 2m decreases from \( N \) toward \( N/2 \).

The kagomé layers of volborthite satisfy the three conditions necessary for a crystalline solid to exhibit a purely entropy-driven phase transitions: [30–32] (1) It has a set of states identical in enthalpy \( H \). (2) The members of this set are grouped into subsets \( i = 1, 2, 3, \ldots \), with degeneracies \( \Omega_1, \Omega_2, \Omega_3, \ldots \), respectively. (3) The degeneracy \( \Omega_i \) increases steadily, e.g., \( \Omega_1 < \Omega_2 < \Omega_3 < \ldots \), so the associated entropy increases steadily. Thus, one may expect that already moderate magnetic fields inducing Zeeman energies comparable to \( J_I \) are able to substantially alter the entropy balance, causing dramatic effects on the low-temperature ordering behavior. In fact, as shown Figure 4a, the specific heat anomalies observed at \( B = 0 \) indeed disappears for \( B > \sim4.2 \) T, but a new anomaly occurs when \( B > \sim5.5 \) T. Below \( \sim2 \) K, neither electronic nor vibrational energy of volborthite is available for the energy absorption, causing the specific heat anomalies. Thus, in the following, we analyze the dependence of the specific heat anomalies on \( T \) and \( B \) in terms of the entropy spectra of Groups I, II and III (Figure 4b).

4.3. Specific Heat Anomalies and Entropy Spectra

As shown above, the entropy \( S_I(2m) \) of Group I is smaller than that of Group II or III by \( S_E(2m) \). As the magnetic field \( B \) increases from 0, a kagomé layer starts absorbing energy by exploiting the entropy spectrum of Group I, so the anomaly below \( \sim4.2 \) T arises from the entropy spectrum of Group I. Since \( \Omega_I(2m) \) becomes larger with increasing 2m from 0 to \( N/2 \) and also with decreasing 2m from \( N \) to \( N/2 \), more states are involved in the energy absorption from Zeeman energy. This allows one to understand why the intensity of the
specific heat anomaly increases with increasing field \( B \). The entropies \( S_I(2m) \) from the states for \( 2m = 0 - N/2 \) as well as \( 2m = N - N/2 \) are used to absorb energy, eventually reaching the highest entropy level \( S_I(N/2) \) (Figure 4b). When all entropy levels are exploited, there is no more entropy level with which a kagomé layer can absorb energy any further, even if \( B \) is increased. At this stage, the spin ladder ordering within a kagomé layer assumes a liquid-like disordered state, and the specific heat in the valley between \(-4.2 \, \text{T} \) and \(-5.5 \, \text{T} \) reaches that of the disordered state of the \( S = 1/2 \) AUH chains (the green plateau in Figure 4a). It is most reasonable to assign the higher-temperature specific heat ridge to a magnetic ordering within each kagomé layer, and the lower-temperature one to a three-dimensional ordering, i.e., an ordering between the magnetically ordered kagomé layers. When the magnetic field increases further, a kagomé layer can utilize the entropy spectra of Groups II and III simultaneously, leading to the single broad specific heat anomaly above \(-5.5 \, \text{T} \). However, this requires that the spin ladder arrangements of a kagomé layer must be converted from Group I to Groups II and III. Each spin ladder has two sets of \( J_1 \) exchanges, e.g., a (AF) spin ladder has one set of AFM-coupled \( J_1 \) and another set of FM-coupled \( J_1 \) exchanges. Suppose that a set of \( J_1 \) paths is switched from FM to AFM, e.g., from an (AF) to a (AA) spin ladder in one member of \( \Omega_{AA}(2m) \) of Group I to create an extra (AA) spin ladder. Such a \( J_1 \)-flip, e.g., \((AF)_m(AF)(AF)_n \rightarrow (AF)_m(FA)(FA)_n\), converts the electronic structure from Group I to Group II, raising the entropy from \( S_I(2m) \) to \( S_{II}(2m) = S_I(2m) + S_L(2m) \). Similarly, a \( J_1 \)-flip can create an extra (FF) spin ladder, e.g., \((FA)_m(FA)(FA)_n \rightarrow (FA)_m(FF)(FF)(FA)_n\), converting the electronic structure from Group I to Group III, raising the entropy from \( S_I(2m) \) to \( S_{III}(2m) = S_I(2m) + S_L(2m) \).

With a magnetic field, it is energetically more favorable to induce a \( J_1 \)-flip from AFM to FM rather than that from FM to AFM. In contrast, thermally-driven \( J_1 \)-flips from AFM to FM or from FM to AFM are equally likely. Thus, thermal agitation will be more effective in creating \( J_1 \)-flips than Zeeman energy, approximately by a factor of 2. The positions of two ridges of the specific heat below \(-4.2 \, \text{T} \) are nearly parallel to the magnetic field axis, whereas the single specific heat ridge above \(-5.5 \, \text{T} \) is slanted with slope \( g\mu_B B/k_B T \approx 2 \). This reflects that, at a strong enough magnetic field, the kagomé layers of Group II spin ladder arrangements absorb energy equally, as do those of Group III spin ladder arrangements.

The entropy spectra II and III are identical in the degeneracy of each entropy level, but differ in the nature of spin ladder arrangements. Whether a given kagomé layer produces the entropy spectrum II or III has no effect on the enthalpy difference. Consequently, there is no ordering between kagomé layers, which explains the occurrence of a single specific heat anomaly at magnetic field above \(-5.5 \, \text{T} \). The latter forms a single ridge with increasing \( B \), the shape of which suggests that not all available entropy states of the entropy spectra II and III are populated at \( 9 \, \text{T} \). If it is assumed that this above-5.5 T specific heat anomaly (Figure 4a) retains a symmetric shape with respect to the temperature and magnetic field, one might expect that the magnetic order above \(-5.5 \, \text{T} \) will disappear under a magnetic field higher than \(-13 \, \text{T} \).

Finally, we note that in all ordered magnetic states of Groups I, II and III, there occurs a strong gain of translational entropy with respect to those states in which there is no order between neighboring spin ladders, because the ordered states have translational symmetry along the leg direction. This provides a strong driving force for the magnetic ordering. In a sense, this situation is similar to the nematic phase transition in a system of thin rods examined by Onsager [31].

5. Concluding Remarks

The spin exchanges of volborthite show that each kagomé layer of Cu\(^{2+}\) ions is hardly spin-frustrated, but rather consists of very weakly interacting two-leg spin ladders with linear trimers as rungs. Below \(-75 \, \text{K} \), these rungs act as \( S = 1/2 \) pseudospin units, making each two-leg spin ladder behave as a \( S = 1/2 \) antiferromagnetic uniform Heisenberg chain. This conclusion was confirmed by synthesizing single crystal samples of volborthite \( \text{Cu}_3\text{V}_2\text{O}_7(\text{OH})_2 \cdot 2\text{H}_2\text{O} \) and its deuterated analogue \( \text{Cu}_3\text{V}_2\text{O}_7(\text{OD})_2 \cdot 2\text{D}_2\text{O} \) and sub-
sequently measuring their magnetic susceptibilities and specific heat anomalies. Under a magnetic field $B$ higher than ~4.2 T, the specific heat anomalies below 1.5 K are suppressed. However, a new specific heat anomaly appears when $B$ is raised above ~5.5 T. The dependence of the specific heat anomalies on the magnetic field are governed by the fact that the magnetic properties of each kagomé has three sets of entropy spectra with which it can exchange energy with the surrounding. These entropy spectra arise from the topologically constrained interactions between adjacent two-leg spin ladders in each kagomé layer; that all spin exchange paths forming each two-leg spin ladder should remain antiferromagnetically coupled.

The present work makes it clear that the use of a correct spin lattice is critical in describing the properties of a magnetic material. As a criterion for finding a proper spin lattice, the geometrical pattern of magnetic ion arrangement can be grossly misleading because a spin lattice is decided by the geometrical pattern of the strong spin exchange paths [3,4,19]. If volborthite were to be treated as a spin-frustrated kagomé lattice model, the $S = 1/2$ antiferromagnetic uniform Heisenberg chain behaviors of its magnetic susceptibility and magnetization are exotic and novel, as are its specific heat anomalies below 1.5 K. By the same token, all other magnetic properties of volborthite not explained by a spin-frustrated kagomé lattice model would be novel and surprising. However, our work showed that each kagomé layer of Cu$^{2+}$ ions consists of very-weakly interacting two-leg spin ladders, so the seemingly exotic magnetic properties previously attributed to magnetic frustration are simply explained by a well-studied $S = 1/2$ antiferromagnetic uniform Heisenberg chain model. To find a correct spin lattice for any given magnetic material, it is necessary to evaluate the relative strengths of various possible spin exchanges of a given magnetic system by using an unbiased and straightforward method such as the energy-mapping analysis, based on first principles DFT calculations [3,4,19,33].

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References
1. Mendels, P.; Wills, A.S. Kagomé antiferromagnets: Materials vs. spin liquid behaviors. In Introduction to Frustrated Magnetism; Lacroix, C., Mendels, P., Mila, F., Eds.; Springer: Berlin/Heidelberg, Germany, 2011; pp. 207–238.
2. Syzranov, S.V.; Ramirez, A.P. Hidden energy scale in geometrically frustrated systems: Are quantum spin liquids achievable? arXiv 2021, arXiv:2105.08070.
3. Xiang, H.J.; Lee, C.; Koo, H.-J.; Gong, X.; Whangbo, M.-H. Magnetic properties and energy-mapping analysis. Dalton Trans. 2013, 42, 823–853. [CrossRef] [PubMed]
4. Whangbo, M.-H.; Xiang, H.J. Magnetic properties from the perspectives of electronic hamiltonian: Spin exchange parameters, spin orientation and spin-half misconception. In Handbook in Solid State Chemistry, Volume 5: Theoretical Descriptions; Dronskowskii, R., Kikkawa, S., Klein, A., Eds.; Wiley: Hoboken, NJ, USA, 2017; pp. 285–343.
5. Hiroi, Z.; Ishikawa, H.; Yoshida, H.; Yamamura, J.; Okamoto, Y. Orbital transitions and frustrated magnetism in the kagome-type copper mineral volborthite. Inorg. Chem. 2019, 58, 11949–11960. [CrossRef]
6. Ishikawa, H.; Yamaura, J.I.; Okamoto, Y.; Yoshida, H.; Nilsen, G.J.; Hiroi, Z. A novel crystal polymorph of volborthite, Cu$_3$V$_2$(OH)$_2$·2H$_2$O. Acta Cryst. C 2012, 68, i41–i44. [CrossRef] [PubMed]
7. Yoshida, H.; Yamamura, J.; Isobe, M.; Okamoto, Y.; Nielson, G.J.; Hiroi, Z. Orbital switching in a frustrated magnet. Nat. Commun. 2012, 29, 860. [CrossRef] [PubMed]
8. Ishikawa, H.; Yoshida, M.; Nawa, K.; Jeong, M.; Krämer, S.; Horvatić, M.; Berthier, C.; Takigawa, M.; Akaki, M.; Miyake, A.; et al. One-third magnetization plateau with a preceding novel phase in volborthite. Phys. Rev. Lett. 2015, 114, 227202. [CrossRef]
1. Nakamura, D.; Yamashita, T.; Ishikawa, H.; Hiroi, Z.; Takeyama, S. Extremely wide 1/3 magnetic plateau of volborthite Cu$_3$V$_2$O$_7$(OH)$_2$·2H$_2$O in ultrahigh magnetic fields up to 182 T. Phys. Rev. B 2018, 98, 020404. [CrossRef]

2. Yamashita, S.; Moriura, T.; Nakazawa, Y.; Yoshida, H.; Okamoto, Y.; Hiroi, Z. Thermodynamic properties of the kagomé lattice in volborthite. J. Phys. Soc. Jpn. 2010, 79, 083710. [CrossRef]

3. Janson, O.; Furukawa, S.; Momoi, T.; Sindzingre, P.; Richter, J.; Held, K. Magnetic behavior of volborthite Cu$_3$V$_2$O$_7$(OH)$_2$·2H$_2$O determined by coupled trimers rather than frustrated chains. Phys. Rev. Lett. 2016, 117, 037206. [CrossRef]

4. Kresse, G.; Furthmüller, J. Efficiency of ab-initio total energy calculations for metals and semiconductors using a plane-wave basis set. Comput. Mater. Sci. 1996, 6, 15–50. [CrossRef]

5. Kresse, G.; Furthmüller, J. Efficient iterative schemes for ab initio total-energy calculations using a plane-wave basis set. Phys. Rev. B 1996, 54, 11169–11186. [CrossRef] [PubMed]

6. Kresse, G.; Joubert, D. From ultrasoft pseudopotentials to the projector augmented-wave method. Phys. Rev. B 1999, 54, 13417–13423. [CrossRef] [PubMed]

7. Blöchl, P.E. Projector augmented-wave method. Phys. Rev. B 1994, 50, 17953–17979. [CrossRef] [PubMed]

8. Perdew, J.P.; Burke, K.; Ernzerhof, M. Generalized gradient approximation made simple. Phys. Rev. Lett. 1996, 77, 3865–3868. [CrossRef]

9. Dudarev, S.L.; Botton, G.A.; Savrasov, S.Y.; Humphreys, C.J.; Sutton, A.P. Electron-energy-loss spectra and the structural stability of nickel oxide: An LSDA+U study. Phys. Rev. B 1998, 57, 1505–1509. [CrossRef]

10. Yan, H.W.; Luo, Y.Z.; Xu, X.; He, L.; Tan, J.; Li, Z.H.; Hong, X.F.; He, P.; Mai, L.Q. Facile and Scalable Synthesis of Zn$_2$V$_2$O$_7$(OH)$_2$·2H$_2$O Microflowers as a High-Performance Anode for Lithium-Ion Batteries. ACS Appl. Mater. Interfaces 2017, 9, 27707–27714. [CrossRef] [PubMed]

11. Whangbo, M.-H.; Koo, H.-J.; Kremer, R.K. Spin exchanges between transition metal ions governed by the ligand p-orbitals in the frustrated Heisenberg chain system cupric chloride CuCl$_2$ (PO$_4$)$_2$. Acta Cryst. C 1979, 55, 501–504. [CrossRef]

12. Selwood, P.W. Magnetochemistry; Interscience Publishers Inc.: New York, NY, USA, 1956.

13. Johnston, D.C.; Kremer, R.K.; Troyer, M.; Wang, X.; Klümper, A.; Bud’ko, L.; Panchula, A.F.; Canfield, P.C. Thermodynamics of spin S=1/2 antiferromagnetic uniform and alternating-exchange Heisenberg chains. Phys. Rev. B 2000, 61, 9588–9606. [CrossRef]

14. B Gwen, P.W. Magnetochemistry; Interscience Publishers Inc.: New York, NY, USA, 1956.

15. Bain, G.A.; Berry, J.F. Diamagnetic corrections and Pascal’s constants. J. Chem. Educ. 2008, 85, 532–536. [CrossRef]

16. Banks, M.G.; Kremer, R.K.; Hoch, C.; Simon, A.; Ouladdiaf, B.; Broto, J.-M.; Rakoto, H.; Lee, C.; Whangbo, M.-H. Magnetic ordering in the frustrated Heisenberg chain system cupric chloride CuCl$_2$. Phys. Rev. B 2009, 80, 024404. [CrossRef]

17. Abraam, A.; Blaney, B. Electron Paramagnetic Resonance of Transition Ions; Oxford University Press: New York, NY, USA, 1970.

18. Poukhari, A.; Moquine, A.; Flandrois, S. A new linear trimeric magnetic ion in (Ca,Sr)$_2$V$_2$O$_7$(OH)$_2$·2H$_2$O phosphates. Mat. Res. Bull. 1986, 21, 395–400. [CrossRef]

19. Todo, S.; Kato, K. Cluster algorithms for general-S quantum spin systems. Phys. Rev. Lett. 2001, 87, 047203. [CrossRef] [PubMed]

20. Albuquerque, A.F.; Alet, F.; Corboz, P.; Dayal, P.; Feiguin, A.; Fuchs, S.; Gamper, L.; Gull, E.; Gürtler, S.; Honecker, A.; et al. The ALPS project release 1.3: Open-source software for strongly correlated systems. J. Magn. Magn. Mater. 2007, 310, 1187–1193. [CrossRef]

21. Zavalij, Y.Y.; Zhano, F.; Whittingham, M.S. A new pyrovannadate, Zn$_3$(OH)$_2$V$_2$O$_7$·2H$_2$O, from X-ray powder data. Acta Cryst. C 1997, 53, 1738–1739. [CrossRef]

22. Boo, W.O.J.; Stout, J.W. Heat capacity and entropy of CuF$_2$ and CrF$_2$ from 10 to 300 K. Anomalies associated with magnetic ordering and evaluation of magnetic contributions to the heat capacity. J. Chem. Phys. 1979, 71, 9–16. [CrossRef]

23. Frenkel, D. Entropy driven phase transitions. Physica A 1999, 263, 26–38. [CrossRef]

24. Karayiannis, N.C.; Foteinopoulou, K.; Laso, M. Entropy-driven crystallization in dense systems of a thermal chain molecules. Phys. Rev. Lett. 2009, 103, 045703. [CrossRef]

25. Orsager, L. The effects of shape on the interaction of colloidal particles. Ann. N. Y. Acad. Sci. 1949, 51, 627–659. [CrossRef]

26. Li, X.; Yu, H.; Lou, F.; Feng, J.; Whangbo, M.-H.; Xiang, H.J. Spin Hamiltonians in magnets: Theories and computations. Molecules 2021, 26, 803. [CrossRef]