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Magnetically ordered and kagome quantum spin liquid states in the Zn-doped claringbullite series

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Neutron scattering measurements have been performed on deuterated powder samples of claringbullite and Zn-doped claringbullite $[\text{Zn}_x\text{Cu}_{4-x}(\text{OD})_6\text{FCl}]$. At low temperatures, claringbullite $\text{Cu}_4(\text{OD})_6\text{FCl}$ forms a distorted pyrochlore lattice with long-range magnetic order and spin-wave-like magnetic excitations. Partial Zn doping leads to the nominal $\text{ZnCu}_3(\text{OD})_6\text{FCl}$ compound, a geometrically frustrated spin-1/2 kagome antiferromagnet that shows no transition to magnetic order down to 1.5 K. The magnetic excitations form a gapless continuum, a signature of fractional excitations in a quantum spin liquid.

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I. INTRODUCTION

Interacting spins forming a two-dimensional lattice of corner sharing triangles, the kagome lattice, are expected to form unconventional magnetic ground states due to geometric frustration of the exchange interactions. These ground states are predicted to include zero-temperature long-range ordered states with noncollinear and even noncoplanar multi- \mathbf{k} magnetic orders for classical spins [1]. For quantum spins, in particular spin 1/2, a large variety of more exotic quantum spin liquid (QSL) ground states have been predicted [2–4]. A crucial question for the classification of a spin liquid is whether the excitation spectrum has a gap or not [5].

Despite much effort, there are relatively few good experimental model systems to study spin-1/2 kagome QSLs. One example is the mineral herbertsmithite, $\text{ZnCu}_3(\text{OH})_6\text{Cl}_2$, where the magnetic response resembles a collection of uncorrelated nearest-neighbor (NN) singlets [6]. The existence of a gap for this state is currently under discussion [7,8], chiefly because of Cu/Zn mixing on the Cu site. Such site disorder has also hampered the study of the α polymorph of herbertsmithite, the mineral kapellasite [9,10], where a competition between NN ferromagnetic $J_1 < 0$ and an antiferromagnetic third-neighbor interaction J_d [9,11] (see Fig. 1 for a definition of the exchanges) leads to a gapless chiral quantum spin liquid within the cuboc2 part of the classical phase diagram [1,3].

In this work, we study another candidate kagome QSL, $\text{ZnCu}_3(\text{OD})_6\text{FCl}$ (hereafter referred to as ZnCu_3), obtained by partial Zn doping of the mineral claringbullite $\text{Cu}_4(\text{OD})_6\text{FCl}$ (hereafter referred to as Cu_4). In the latter, three of the four Cu^{2+} ions in the unit cell form perfect spin-1/2 kagome layers at room temperature. The fourth Cu^{2+} ion sits alternatively

above and below the kagome triangles, leading to a slightly distorted pyrochlore lattice. The resulting distorted trigonal prism coordination of the interplanar Cu^{2+} ions contrasts with the distorted octahedral coordination of herbertsmithite. This coordination geometry is expected to drastically reduce Cu/Zn site disorder in ZnCu_3 , since the closed-shell Zn^{2+} ion prefers the interlayer site compared to the more distorted octahedral sites of the kagome plane, as predicted in the isostructural mineral barlowite $\text{Cu}_4(\text{OH})_6\text{FBr}$ [12].

Claringbullite (Cu_4), first reported with a slightly different chemical formula [13], crystallizes at room temperature in the hexagonal space group $P6_3/mmc$ (No. 194) [14]. The interlayer Cu, which is disordered over three equivalent positions in this structure, orders slightly below room temperature, leading to an orthorhombic distortion and a doubling of the crystallographic unit cell [15,16]. The magnetic interactions in claringbullite are predominantly antiferromagnetic, as seen from the Weiss temperature $\theta_W = -135$ K, and magnetic order is established below $T_N = 17$ K [16–18]. Zinc-doped claringbullite (ZnCu_3), on the other hand, remains in the $P6_3/mmc$ space group down to the lowest temperatures, as will be shown below. It does not order magnetically down to at least $T = 0.8$ K according to specific heat measurements despite a largely antiferromagnetic θ_W of -223 K [18], and is therefore a promising kagome QSL candidate. Here, we present inelastic neutron scattering measurements on highly deuterated $\text{Cu}_4(\text{OD})_6\text{FCl}$ and $\text{ZnCu}_3(\text{OD})_6\text{FCl}$ powder samples, and their crystallographic and magnetic characterization.

II. SYNTHESIS AND CHARACTERIZATION

A. Synthesis

Highly deuterated Cu_4 and ZnCu_3 polycrystalline samples of weight 5–7 g were synthesized using a hydrothermal

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method adapted from [18], as detailed in the Supplemental Material [19].

B. Crystallographic studies

Laboratory x-ray powder diffraction ($\lambda = 1.5406$ Å) showed both the Cu_4 and ZnCu_3 samples to be single phase and to crystallize in the $P6_3/mmc$ space group at room temperature. This was confirmed by high-resolution powder neutron diffraction using D2B at the Institut Laue-Langevin (ILL), where measurements were performed with $\lambda = 1.5952$ Å at $T = 1.5$ and 295 K [20]. Rietveld refinements showed a high deuteration level of $\sim 96\%$.

At $T = 1.5$ K, the powder neutron diffraction data from Cu_4 show additional peaks that were successfully indexed in the $Pnma$ space group [$a = 11.5359(9)$, $b = 9.1510(7)$, and $c = 6.6848(5)$ Å], in agreement with previous single crystal studies at $T = 100$ K [15]. The driving force for this distortion is the ordering of the interlayer Cu onto one of three local equivalent sites. At low temperatures, the Cu_4 $Pnma$ structure is a distorted pyrochlore with three Cu sites at the $4a$, $8d$, and $4c$ Wyckoff positions. The kagome to interlayer Cu distances range from 2.80 to 3.17 Å, slightly smaller than the kagome Cu-Cu distances that range from 3.32 to 3.35 Å.

ZnCu_3 does not show additional peaks at $T = 1.5$ K, indicating that it remains in the $P6_3/mmc$ space group [$a = 6.65918(6)$ and $c = 9.17288(9)$ Å]. The refinement of the low temperature neutron diffraction data shows that the kagome site is within the error of the refinement fully occupied by Cu. The interlayer position is occupied by $\sim 74\%$ Zn on the high-symmetry $2d$ site while the Jahn-Teller active Cu ion occupies $\sim 8.5\%$ of the $6h$ site (see Supplemental Material [19] and references [21,22] therein).

C. Magnetometry

Magnetic susceptibility measurements in a magnetic field of 1000 G showed a magnetic transition in Cu_4 at $T_N \approx 17$ K with a weak ferromagnetic component (possibly due to Dzyaloshinskii-Moriya interactions) and a Weiss temperature of $\theta_W = -136(3)$ K (see Supplemental Material [19]). In ZnCu_3 , no transition to magnetic order was seen down to 2 K and the Weiss temperature was $\theta_W = -206(1)$ K (Supplemental Fig. S5 [19]), in agreement with [18].

Magnetization measurements as a function of magnetic field show, contrary to the literature [18], a small hysteresis in ZnCu_3 of 0.06 T with a spontaneous moment of $2 \times 10^{-4} \mu_B/\text{Cu}$ at $T = 2$ K, which corresponds to 0.01% of the full ordered moment (see Supplemental Fig. S5 [19]). This could be intrinsic to our ZnCu_3 sample; however, we note that the hysteresis loop opens below 6 K, which corresponds to the ordering temperature of clinoatacamite [23]. Although no impurity phase was observed in the D2B neutron diffraction data, previous work has reported impurities of polymorphs of clinoatacamite [16] or herbertsmithite [24] in both Cu_4 and ZnCu_3 . The observed spontaneous moment at 2 K could be attributed to a 0.4% clinoatacamite impurity [25], which would not be visible in our powder diffraction data.

III. MAGNETIC STRUCTURE

The magnetic structure of Cu_4 was determined at $T = 1.7$ K using the high-flux neutron diffractometer D20 at the

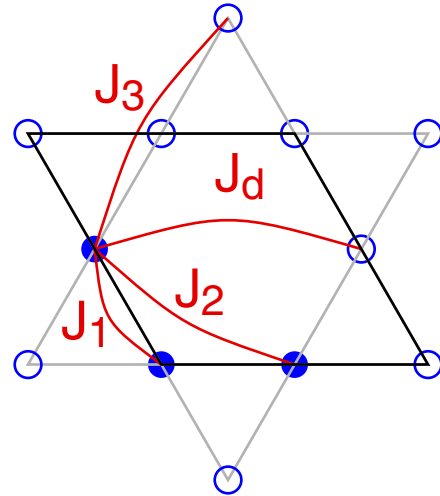


FIG. 1. Kagome lattice showing the first four exchange interactions. Black lines show the unit cell with three Cu^{2+} $S = 1/2$ ions as filled circles.

ILL with $\lambda = 2.410$ Å [26]. Subtraction of data taken at $T = 25$ K, just above the magnetic phase transition at $T_N = 17$ K, show more than a dozen magnetic peaks that correspond to an antiferromagnetic order with propagation vector $\mathbf{k} = 0$ in the orthorhombic space group (see Supplemental Material [19]). Refinements of the different irreducible representations (IRs) obtained by representation analysis [27] gave a good fit only for the IR Γ_7 , in Kovalev's notation, which corresponds to the magnetic space group $Pn'm'a$, previously found for both barlowite [16,28] and claringbullite [16]. The ordered magnetic moments are found to be reduced from the fully ordered $1\mu_B$ expected for $S = 1/2$, which we relate to quantum fluctuations, with values of $0.26\mu_B$, $0.37\mu_B$, and $0.52\mu_B$ for the Cu1 site at $(0, 0, 0)$, Cu2 at $(0.247, 0.496, 0.249)$, and Cu3 at $(0.191, 1/4, 0.051)$, respectively. Details of the basis vectors for the IR Γ_7 can be found in the Supplemental Material [19].

The magnetic moments are mostly aligned along the orthorhombic a axis, with the Cu2 moment rotated (antiferromagnetically) by 18° towards b and the Cu3 moment canted ferromagnetically by 17° towards c ; see Fig. 2(a). Of the three spins on each kagome triangle, two are almost parallel to the a axis, and so to the interlayer spin, and one is antiparallel. The former are closer in distance to the interlayer spin, and one may speculate that the “tripod” interactions J_{tn} ($n = 1, 2, 3$) between the interlayer and the kagome spins are ferromagnetic, in agreement with density functional theory calculations for barlowite [29]. This scenario is also supported by the smaller negative value of the Weiss temperature of Cu_4 compared to ZnCu_3 , which suggests additional ferromagnetic interactions in the former. The canting of the spins away from the a axis indicates that antisymmetric Dzyaloshinskii-Moriya (DM) interactions may play a role. Measurements at $T = 15$ K, just slightly below T_N , show the same magnetic structure as at $T = 1.7$ K, with further reduced moment sizes due to thermal fluctuations.

The magnetic structure found for Cu_4 in this work is similar to that reported for a different polycrystalline sample [16],

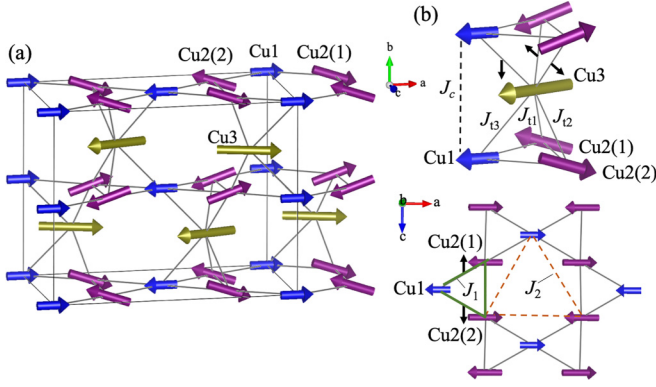


FIG. 2. (a) Magnetic structure of $\text{Cu}_4(\text{OD})_6\text{FCl}$ at $T = 1.7$ K. (b) Relevant exchange interactions in the orthorhombic structure. The black arrows indicate the Dzyaloshinskii-Moriya vectors along the [010] and [001] directions (see Table I).

with the main difference being 20–30% smaller moment sizes in our samples. The magnetic structure also resembles that of barlowite, which has slightly larger canting angles: $\sim 28^\circ$ for Cu2 and $\sim 22^\circ$ for Cu3 [28].

IV. INELASTIC NEUTRON SCATTERING

A. Measurements

Inelastic neutron scattering (INS) measurements on ZnCu_3 were performed on the cold-neutron time-of-flight (TOF) spectrometer LET (ISIS) using neutrons with incoming energies of $E_i = 2.8, 6.0$, and 20 meV at temperatures of 1.7 and 50 K [30], while Cu_4 was measured on the cold and thermal TOF spectrometers IN5 and Panther (ILL) using incoming neutron energies between 3.5 and 35 meV at temperatures between 1.7 and 100 K [31,32]. Standard data reduction was carried out using MANTID [33]. For comparison between the two samples, the scattering was put on an absolute scale by normalizing to the intensities of nuclear Bragg peaks.

B. Zn-doped claringbullite $\text{ZnCu}_3(\text{OD})_6\text{FCl}$

No magnetic Bragg peaks were observed in data collected for ZnCu_3 at $T = 1.7$ K, which confirms the absence of magnetic order in this compound. The scattering function $S(Q, E)$ at $T = 1.7$ K shows a low-energy spin-liquid-like magnetic response centered at about $Q = 0.7 \text{ \AA}^{-1}$ with an energy-independent full width at half maximum in Q of about 0.7 \AA^{-1} ; see Fig. 3(a). The excitations are peaked at about 1 meV with a non-Lorentzian energy profile and a band width of about 3 meV. The excitations have no discernible energy gap, at least down to 0.27 meV.

In addition to the clear low-energy magnetic response, there is a second contribution at higher energies that also appears to be magnetic. It is centered at $Q \approx 1.2 \text{ \AA}^{-1}$ [see Fig. 3(b)] and has a typical width of about 0.7 \AA^{-1} , which corresponds to a correlation length of 8.4 \AA (approximately 1.2 kagome hexagons). This excitation has a band width of about 10 meV and no discernible gap.

These two magnetic responses have different energy ranges but can both be indexed by a $\mathbf{k} = \mathbf{0}$ characteristic wave vector,

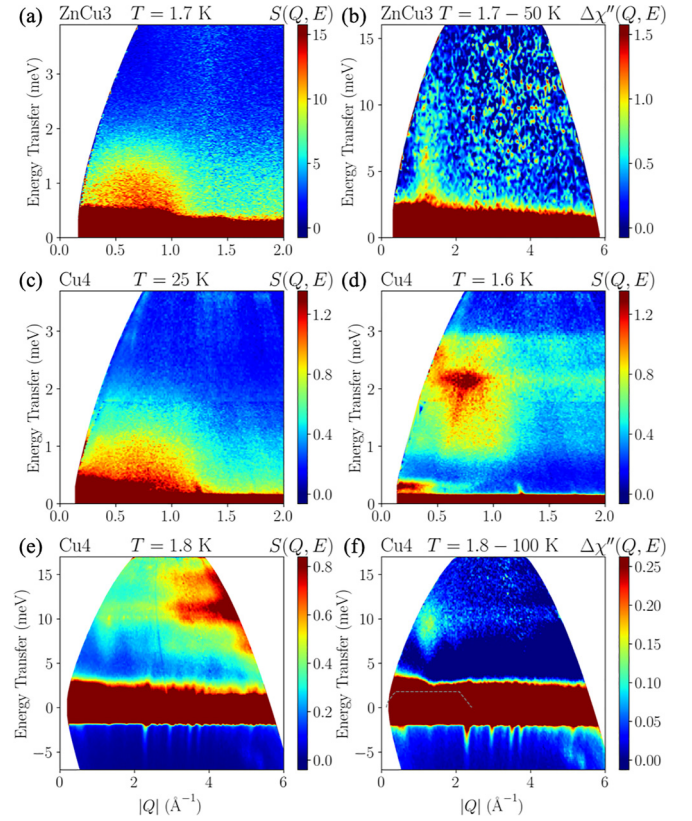


FIG. 3. (a)–(b) INS spectra from $\text{ZnCu}_3(\text{OD})_6\text{FCl}$ at $T = 1.7$ K measured on LET: (a) $S(Q, E)$ for $E_i = 6.0$ meV; (b) Difference in magnetic dynamic susceptibility $\chi''(Q, E)$ between $T = 1.7$ and 50 K for $E_i = 20$ meV. (c)–(d) $S(Q, E)$ of $\text{Cu}_4(\text{OD})_6\text{FCl}$ at two different temperatures measured on IN5 by combining $E_i = 3.55$ and 5.1 meV: (c) Cooperative paramagnetic state at $T = 25$ K; (d) Magnetically ordered state at $T = 1.6$ K. (e)–(f) INS spectra from $\text{Cu}_4(\text{OD})_6\text{FCl}$ measured on Panther with $E_i = 19$ meV: (e) $S(Q, E)$ at $T = 1.8$ K; (f) Difference in magnetic dynamic susceptibility $\chi''(Q, E)$ between $T = 1.8$ and 100 K showing both the low-energy response in the tail of the elastic peak and the high-energy response near 10 meV. All data in this figure are in arbitrary units.

corresponding to $(0,0,1)$ and $(1,0,1)$ reflections, respectively. At $T = 50$ K, both magnetic responses persist but are weaker in intensity.

C. Claringbullite $\text{Cu}_4(\text{OD})_6\text{FCl}$

In Cu_4 above the ordering temperature, a low-energy magnetic response is seen at $Q = 0.7 \text{ \AA}^{-1}$ [see Fig. 3(c)], very similar to that observed in ZnCu_3 . In the magnetically ordered phase, this scattering develops features strongly resembling dispersive spin waves, see Fig. 3(d), with an energy gap of 0.45 meV and a band width of 3.35 meV. Additionally, there is a high-energy magnetic response centered at $Q \approx 1.2 \text{ \AA}^{-1}$, with a 5 meV energy gap and scattering extending up to 13 meV [see Figs. 3(e) and 3(f)].

V. ANALYSIS

A. Spin waves in $\text{Cu}_4(\text{OD})_6\text{FCl}$

An attempt was made to describe the observed spin waves of Cu_4 using semiclassical linear spin wave theory with the

TABLE I. Exchange interactions used for the spin-wave calculations. The labels refer to Fig. 2(b) and positive values denote antiferromagnetic couplings. Dzyaloshinskii-Moriya interactions along the [010] and [001] directions are given.

Label	J_i (meV)	$ D_{[010]} /J_i$	$ D_{[001]} /J_i$
J_1	14.0	0	0.03
J_{t1}	-6.5	0	0.03
J_{t2}	-3.5	0	0.03
J_{t3}	-5.9	0.05	0
J_c	4.0	0	0
J_2	0.2	0	0

spin Hamiltonian

$$\mathcal{H} = \sum_{i,j} J_{ij} \mathbf{S}_i \cdot \mathbf{S}_j + \mathbf{D}_{ij} \sum_{i,j} \mathbf{S}_i \times \mathbf{S}_j. \quad (1)$$

It turned out to be quite difficult to stabilize the observed magnetic structure, even when including isotropic exchange interactions J_{ij} up to six further neighbors as well as antisymmetric anisotropic DM interaction terms \mathbf{D}_{ij} . The exchange interactions predicted from combined density functional theory calculations and magnetic susceptibility measurements for barlowite [29], which has crystallographic and magnetic structures similar to claringbullite, also did not stabilize the observed magnetic structure in claringbullite.

The complicated magnetic structure in Cu_4 suggests that several exchange interactions are involved. To explore the sizable phase space of Eq. (1), the program SERENDIPITY [34] was developed to identify cluster areas of phase space where the experimental magnetic structure could be stabilized by a given set of exchange parameters. The magnetic excitation spectra from the resulting sets of exchange integrals were calculated using SPINW [35], allowing for a relaxation of the observed canting angles in the magnetic structure to improve stability, and compared to the experimental data.

The final model includes the exchange interactions J_{ij} shown in Fig. 2(b) and specified in Table I, namely, a strong antiferromagnetic nearest-neighbor interaction J_1 in the kagome plane, three different ferromagnetic “tripod” interactions J_{t1} , J_{t2} , and J_{t3} between the kagome spins and the Cu_3 spin that caps the triangles, an antiferromagnetic interaction J_c between the kagome planes, and a very weak antiferromagnetic next-nearest interaction J_2 in the kagome plane. This set of exchange interactions give a Weiss temperature of $\theta_W = -95$ K, in quite close agreement with the experimental value of $\theta_W = -136$ K, and the signs of the interactions are compatible with the expectations from the superexchange bond angles. DM interactions (with both in- and out-of-plane components) were added to the J_1 , J_{t1} , and J_{t2} exchange paths to stabilize the spin canting of the magnetic structure. The calculated spin wave spectra, shown in the Supplemental Material [19], indicate that the model correctly reproduces the bandwidth of the observed spin waves, including the zero-energy gap and the gap between the upper and lower branches. The intensity ratio between these two branches is also reproduced, as well as the overall Q dependence, but the details of the excitation spectra differ

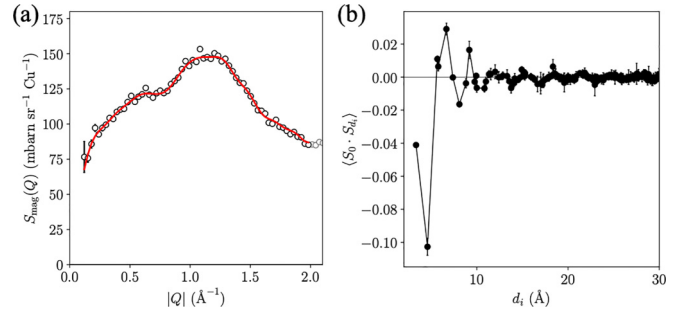


FIG. 4. (a) $S_{\text{mag}}(Q)$ of $\text{ZnCu}_3(\text{OD})_6\text{FCl}$ at $T = 1.7$ K from measurements on LET with $E_i = 2.8, 6.0$, and 20 meV (black symbols) with RMC fit (red line). (b) Radial spin correlations from RMC fits as a function of Cu-Cu distance d .

substantially. Further progress would require measurements on large single crystals, currently unavailable.

B. Zeroth moment analysis of $\text{ZnCu}_3(\text{OD})_6\text{FCl}$

Qualitative information on spin correlations can be obtained using zeroth moment analysis of the neutron scattering intensity $S(Q, E)$. This method was applied to ZnCu_3 data taken at $T = 1.7$ K with incoming energies of $2.8, 6.0$, and 20 meV. The zeroth moment of the magnetic scattering, $S_{\text{mag}}(Q)$, was obtained by integrating the measured (and normalized) $S(Q, E)$ over the energy range $0.1 < E < 12$ meV. To reduce the influence of coherent phonon scattering, which is important for wave vectors $Q > 2 \text{ \AA}^{-1}$, the wave-vector range was limited to below that value, and the data were fed into the SPINVERT program [36], which uses reverse Monte Carlo (RMC) to extract radial spin correlations $\langle \mathbf{S}_0 \cdot \mathbf{S}_d \rangle$. The data of $S_{\text{mag}}(Q)$ and the RMC fit are shown in Fig. 4(a). The refined effective magnetic moment, $\mu_{\text{eff}} = 1.705(1) \mu_B$, is in good agreement with the spin-only value of $1.73 \mu_B$. The radial spin correlations are shown as a function of Cu-Cu distance d in Fig. 4(b). The strongest correlation is negative and corresponds to the interkagome distance, $\sim 4.56 \text{ \AA}$, indicating predominantly antiparallel spin alignment between kagome layers. The nearest-neighbor correlation in the kagome plane ($d \approx 3.33 \text{ \AA}$) is also antiferromagnetic, as expected from the large negative Weiss temperature.

C. Comparison between Cu_4 and ZnCu_3

In the cooperative paramagnetic regime, the magnetic scattering shows very strong resemblances between ZnCu_3 at low temperature and Cu_4 above T_N , as shown in Figs. 3(a) and 3(c). This is further illustrated in Fig. 5 by cuts of $\chi''(Q, E)$ in both constant energy and Q . Scans at constant energy were fitted with three contributions: A Gaussian describing magnetic correlations, a Q -independent term describing uncorrelated magnetic scattering, and an incoherent phonon term given by $a + bQ^2 \exp(-u^2 Q^2/2)$, where a is due to multiple scattering.

At energies of about $5\text{--}6$ meV, Fig. 5(a), the correlated magnetic scattering is centered at $Q \approx 1.2 \text{ \AA}^{-1}$ in both samples, with similar widths in Q , indicating similar correlations in both materials. The scattering underneath these correlations

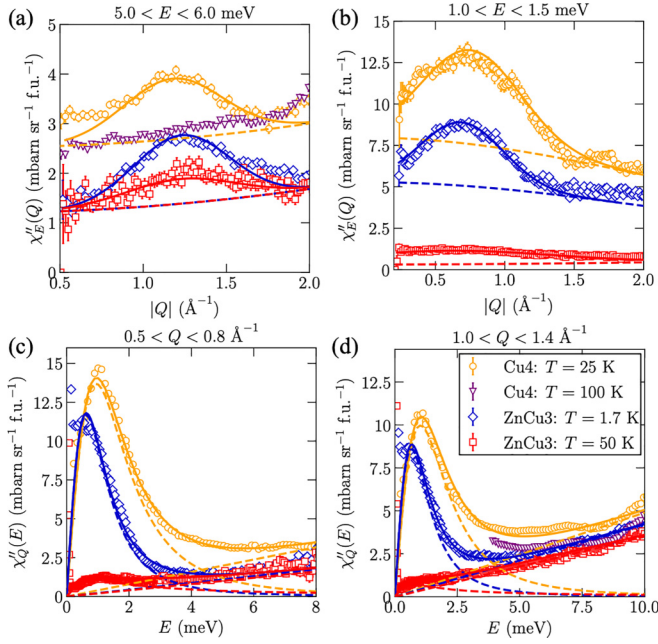


FIG. 5. Comparison of scans in Q [(a)–(b)] and energy [(c)–(d)] of the dynamic magnetic susceptibility in different energy and Q ranges, respectively, of $\text{ZnCu}_3(\text{OD})_6\text{FCl}$ at $T = 1.7$ K (blue) and 50 K (red) and of $\text{Cu}_4(\text{OD})_6\text{FCl}$ at $T = 25$ K (orange) and $T = 100$ K (purple) showing the similarities between the two systems. The solid lines are fits as described in the text. The data are normalized to the number of formula units.

is dominated by incoherent phonons. This phonon scattering is higher in the Cu_4 sample, possibly due to a lower deuteration level, corresponding to 90% D instead of the 96% obtained from the refinement of the diffraction data. The peak areas of Cu_4 at $T = 25$ K and ZnCu_3 at $T = 1.7$ K are within 10% of each other, suggesting that the number of Cu atoms contributing to the higher energy response is the same. This gives strong evidence for the magnetic scattering near $Q = 1.2 \text{ \AA}^{-1}$ being due to correlations from the kagome layers. These correlations persist up to at least $T = 50$ K and possibly vanish at $T = 100$ K.

At lower energies of ~ 1 – 1.5 meV, Fig. 5(b), the correlated magnetic scattering is centered at $Q \approx 0.7 \text{ \AA}^{-1}$ in both samples, with similar widths in Q , again indicating similar correlations in both materials. The broad scattering beneath the correlations is predominantly attributed to uncorrelated magnetic scattering and the remainder arises from incoherent phonons. Both the correlated and uncorrelated magnetic contributions are about 1.5 times stronger in Cu_4 than in ZnCu_3 . This suggests that it is the same atoms that contribute to the correlated and uncorrelated magnetic scattering, and that the scattering comes from all Cu atoms in each sample.

The analysis of neutron diffraction data collected from ZnCu_3 indicates $\sim 8.5\%$ occupancy of the interlayer $6h$ sites by Cu^{2+} . This occupation by a moment bearing ion would introduce some local “tripod” exchange. Comparison of the Q dependences of the magnetic scattering of Cu_4 ($T = 25$ K) and ZnCu_3 ($T = 1.7$ K) shown in Fig. 5 suggests that in this regime the “tripod” exchange in these materials does not notably change the magnetic excitations.

The energy dependence of the excitations was compared by integrating $\chi''(Q, E)$ over wave vector ranges $0.5 < Q < 0.8 \text{ \AA}^{-1}$ and $1.0 < Q < 1.4 \text{ \AA}^{-1}$, shown in Figs. 5(c) and 5(d), respectively. $\chi''(E)$ is peaked at an energy of about 1 meV, and describing it with a power law is therefore not adequate. A quasielastic Lorentzian, often used to describe paramagnetic spins, does not provide a good fit either. This gives further evidence that Zn-doped claringbullite is a quantum spin liquid. We found that a good empirical parametrization of the continuum scattering in ZnCu_3 was achieved using a squared Lorentzian function, $\chi''(E) = 2E\chi'\Gamma^3/[(E^2 + \Gamma^2)^2]$, describing an overdamped excitation where Γ is related to the width in energy of the scattering. An additional contribution, linear in energy, $\chi''(E) = eE$, had to be added to the fits to describe the high-energy behavior. This could arise from both magnetic and phonon scattering, but its very weak temperature dependence indicates that it mainly arises from the density of states of acoustic phonons with energies below the Debye temperature.

At low Q , Fig. 5(c), Cu_4 at $T = 25$ K has a maximum at $E = \Gamma/\sqrt{3} = 0.940(3)$ meV, whereas ZnCu_3 peaks at $0.624(2)$ meV at $T = 1.7$ K. At all Q values, the low-energy magnetic response of Cu_4 has a larger Γ than ZnCu_3 , with values of $1.628(5)$ and $1.081(3)$ meV, respectively, from Fig. 5(c).

VI. DISCUSSION

The model-free zeroth-moment analysis of ZnCu_3 indicates that the strongest correlations are antiferromagnetic and between the kagome layers. Therefore it is likely that these are the main contribution to the low-energy magnetic scattering in ZnCu_3 , as also supported by the observation that the scattering in Fig. 5(b) is centered at $Q = 0.7 \text{ \AA}^{-1}$, corresponding to the $(0,0,1)$ reflection of the $\mathbf{k} = \mathbf{0}$ wave vector in reciprocal space. The low-energy excitation of Cu_4 and ZnCu_3 have very similar profiles with Cu_4 showing additional intensity compared to ZnCu_3 in both the correlated and uncorrelated contributions [see Fig. 5(b)], as well as a larger energy range [see Fig. 5(c)]. This suggests that the additional “tripod” exchange paths introduced by the interlayer Cu atoms in Cu_4 do not significantly change the excitations; instead they strengthen the interkagome correlations present in ZnCu_3 .

These interkagome spin correlations are an important aspect of the Zn-claringbullite family and are likely due to the interkagome distance being shorter than the second nearest-neighbor one in the kagome planes. Exploring possible exchange models for Cu_4 showed that a sizable antiferromagnetic interkagome exchange J_c up to the order of 30% of J_1 , as well as a small further neighbor exchange J_2 , are required to stabilize the magnetic structure. Although the sign of J_c agrees with that previously suggested for barlowite [29], we found that its magnitude relative to J_1 is greater. In ZnCu_3 , the Q positions of the magnetic responses could correspond to the $[001]$ and $[101]$ directions, and hence allude to interkagome spin correlations, shown to be the strongest by the zeroth-moment analysis. Despite the interkagome correlations in ZnCu_3 favoring a three- rather than a two-dimensional system, a previous theoretical study using a coupled cluster method has shown that the kagome quantum spin liquid state can persist up to $J_c/J_1 \approx 15\%$, and it would be

interesting to determine whether ZnCu_3 fits into this QSL phase diagram [37].

In comparison to ZnCu_3 , the powder and single crystal INS data of herbertsmithite also show two magnetic responses centered at $Q \approx 0.67 \text{ \AA}^{-1}$ and $Q \approx 1.3 \text{ \AA}^{-1}$, which extend up to $\sim 2 \text{ meV}$ [6] and $\sim 25 \text{ meV}$ [38], respectively. Although part of the low-energy magnetic scattering was initially attributed to 15% Cu impurity spins on the interlayer sites [39], a variational Monte Carlo study showed it to be an intrinsic kagome response within the nearest-neighbor RVB model [40]. In our analysis of ZnCu_3 , both magnetic responses are attributed to the kagome layers through comparison with the parent material Cu_4 . However, the zeroth-moment analysis indicates that the correlations in ZnCu_3 extend beyond the nearest-neighbor limit, suggesting a different flavor of QSL compared to herbertsmithite. This is further underlined by the difference in energy responses between the two systems: in herbertsmithite the excitations are underdamped [41] while in ZnCu_3 we find them to be overdamped.

The magnetic excitations of both Cu_4 and ZnCu_3 resemble those of the isostructural barlowite and its Zn-doped variant, $\text{Zn}_x\text{Cu}_{4-x}(\text{OD})_6\text{FBr}$ for $0 \leq x < 1$ [42]. For the $x \approx 1$ sample, a finite sized gap of $\sim 0.65 \text{ meV}$ is observed using ^{19}F NMR [43] and inelastic neutron scattering [44]. In contrast, our INS data show ZnCu_3 claringbullite to be gapless down to the lowest resolved energy transfer of $\sim 0.27 \text{ meV}$, but future NMR measurements are needed to verify this.

VII. CONCLUSIONS

Our inelastic neutron scattering measurements show gapless ($\Delta < 0.27 \text{ meV}$), diffuse excitations arising from short-range magnetic correlations in ZnCu_3 down to $T = 1.7 \text{ K}$, suggesting the existence of a quantum spin liquid ground

state. The analysis of the Q and energy dependence of these correlations suggest a different ground state to that of herbertsmithite, widening the library of QSL types in experimental materials. ZnCu_3 shows two inelastic magnetic responses that strongly resemble those of the pyrochlore-like Cu_4 above its magnetic ordering temperature. In contrast, below T_N , the two responses in Cu_4 become gapped from each other and acquire a $\sim 0.5 \text{ meV}$ zero-energy gap. Both responses are attributed to the kagome layers, with the low-energy one arising from interkagome correlations strengthened by the additional interlayer Cu in the parent material via the “tripod” interactions. Single crystal studies will be important in providing a more accurate exchange model for Cu_4 . Local probe measurements, such as NMR or muon spin relaxation studies, may also be helpful in determining whether the ZnCu_3 state remains dynamic to mK temperatures. Furthermore, elemental analysis using anomalous x-ray diffraction could provide more accurate information on the extent of antisite disorder in ZnCu_3 .

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