



THE UNIVERSITY *of* EDINBURGH

Edinburgh Research Explorer

Metamagnetism and soliton excitations in the modulated ferromagnetic Ising chain CoV₂O₆

Citation for published version:

Kimber, SAJ, Mutka, H, Chatterji, T, Hofmann, T, Henry, PF, Bordallo, HN, Argyriou, DN & Attfield, JP 2011, 'Metamagnetism and soliton excitations in the modulated ferromagnetic Ising chain CoV₂O₆' Physical Review B, vol. 84, no. 10, 104425. DOI: 10.1103/PhysRevB.84.104425

Digital Object Identifier (DOI):

[10.1103/PhysRevB.84.104425](https://doi.org/10.1103/PhysRevB.84.104425)

Link:

[Link to publication record in Edinburgh Research Explorer](#)

Document Version:

Publisher's PDF, also known as Version of record

Published In:

Physical Review B

Publisher Rights Statement:

Copyright © 2011 by the American Physical Society. This article may be downloaded for personal use only. Any other use requires prior permission of the author(s) and the American Physical Society.

General rights

Copyright for the publications made accessible via the Edinburgh Research Explorer is retained by the author(s) and / or other copyright owners and it is a condition of accessing these publications that users recognise and abide by the legal requirements associated with these rights.

Take down policy

The University of Edinburgh has made every reasonable effort to ensure that Edinburgh Research Explorer content complies with UK legislation. If you believe that the public display of this file breaches copyright please contact openaccess@ed.ac.uk providing details, and we will remove access to the work immediately and investigate your claim.



Metamagnetism and soliton excitations in the modulated ferromagnetic Ising chain CoV_2O_6 Simon A. J. Kimber,^{1,2,3,*} Hannu Mutka,⁴ Tapan Chatterji,⁴ Tommy Hofmann,² Paul. F. Henry,^{2,5} Heloisa N. Bordallo,^{2,5,6} Dimitri N. Argyriou,^{2,5} and J. Paul Attfield^{3,†}¹European Synchrotron Radiation Facility (ESRF), 6 rue Jules Horowitz, Boîte Postale 220, F-38043 Grenoble Cedex 9, France²Helmholtz-Zentrum Berlin für Materialien und Energie (HZB), Hahn-Meitner Platz 1, D-14109 Berlin, Germany³School of Chemistry and Centre for Science at Extreme Conditions, King's Buildings, West Mains Road, Edinburgh, EH9 3JJ, United Kingdom⁴Institut Max von Laue-Paul Langevin, 6 rue Jules Horowitz, Boîte Postale 156, F-38042 Grenoble Cedex 9, France⁵European Spallation Source ESS AB, Box 176, S-22100 Lund, Sweden⁶Niels Bohr Institute, University of Copenhagen, Universitetsparken 5, 2100 Copenhagen, Denmark

(Received 26 May 2011; revised manuscript received 27 July 2011; published 15 September 2011)

We report a combination of physical property and neutron scattering measurements for polycrystalline samples of the one-dimensional spin-chain compound CoV_2O_6 . Heat capacity measurements show that an effective $S = 1/2$ state is found at low temperatures and that magnetic fluctuations persist up to $\sim 6T_N$. Above $T_N = 6.3$ K, measurements of the magnetic susceptibility as a function of T and H show that the nearest-neighbor exchange is ferromagnetic. In the ordered state, we have discovered a crossover from a metamagnet with strong fluctuations between 5 K and T_N to a state with a $1/3$ magnetization plateau at $2 < T < 5$ K. We use neutron powder diffraction measurements to show that the antiferromagnetic state has incommensurate long-range order and inelastic time-of-flight neutron scattering to examine the magnetic fluctuations as a function of temperature. Above T_N , we find two broad bands between 3.5 and 5 meV and thermally activated low-energy features which correspond to transitions within these bands. These features show that the excitations are deconfined solitons rather than the static spin reversals predicted for a uniform ferromagnetic Ising spin chain. Below T_N , we find a ladder of states due to the confining effect of the internal field. A region of weak confinement below T_N , but above 5 K, is identified which may correspond to a crossover between two- and three-dimensional magnetic ordering.

DOI: [10.1103/PhysRevB.84.104425](https://doi.org/10.1103/PhysRevB.84.104425)

PACS number(s): 75.10.Pq, 78.70.Nx

I. INTRODUCTION

One-dimensional magnetic materials with quantum ($S = 1/2$) spins have been widely studied as their excitation spectra often consist of continuums of fractionalized particles^{1,2} which have no analogs in higher spin or three-dimensional materials. A particularly elegant example is the Ising spin chain, which in a transverse field undergoes one of the best understood quantum phase transitions.^{3–5} Much of the interesting physics in Ising chains, such as propagation of domain walls (solitons), only occurs when neighboring chains are weakly coupled together in real materials. For example, the excitations of an isolated antiferromagnetic Ising chain consist of blocks of reversed spins. Intrachain coupling mixes these states together, creating a broad band centered in energy at twice the interchain coupling constant.⁶ Thermal population of these states creates a low-energy response (the Villain mode), which has a characteristic square-root divergence and can be observed by inelastic neutron scattering.

Recently, attention has focused on those materials which have structures more complicated than simple linear chains. The zigzag chain material CoNb_2O_6 is a notable example, showing a transition from spin-density wave order to noncollinear magnetic order on cooling.^{7,8} Measurements of physical properties^{9,10} and neutron scattering^{11–15} under magnetic fields also detect a rich phase diagram which includes regions of three-phase coexistence.¹⁶

Further increases in structural complexity can also produce ground states which are highly sensitive to external perturbations such as magnetic fields. For example, $\text{Ca}_3\text{Co}_2\text{O}_6$ has a structure consisting of CoO_6 chains on a triangular lattice¹⁷ and undergoes a counterintuitive magnetic *disordering* at

the lowest temperatures. Applied fields in this material find a cascade of magnetization plateaus,¹⁸ which has been attributed to quantum tunneling effects similar to those seen in molecular magnets.¹⁹

In this paper we report our observations on CoV_2O_6 , which crystallizes in two polymorphs.^{20,21} Here we study the triclinic ($P\bar{1}$) brannerite polymorph. (The basic properties of the monoclinic polymorph have also recently been reported.²²) The structure of CoV_2O_6 [Fig. 1(a)] consists of well-separated edge-sharing chains of CoO_6 octahedra running down the b axis interspersed with nonmagnetic V^{5+} in V_2O_5 -like blocks.²¹ The Co-Co distance within the chains is ~ 3 Å and the distances between chains are ~ 4.8 Å along [001] and ~ 7.2 Å along [100]. The chains contain two crystallographically independent sites [Fig. 1(b)], which alternate in the sequence 1, 1, 2, 1, 1, etc. The main magnetic exchange interactions are also shown in Fig. 1(b), with a notable modulation along the chain direction (i.e., $J_1 \neq J'_1$). The magnetic properties of several other brannerites have been reported. The isostructural material CuV_2O_6 is a low-dimensional antiferromagnet, due to Cu^{2+} orbital order,²³ whereas monoclinic MnV_2O_6 is an isotropic antiferromagnet that shows reduced magnetic coherence lengths due to Mn-V antisite disorder.²⁴

II. EXPERIMENT

Polycrystalline samples of CoV_2O_6 were synthesized using a citrate decomposition method²⁴ similar to that used for MnV_2O_6 . Stoichiometric quantities of cobalt(II) acetate tetrahydrate (Aldrich, 99%+) and V_2O_5 (Aldrich, 99.99%) were mixed in distilled water together with a threefold molar excess of acetic acid. The mixture was slowly heated and

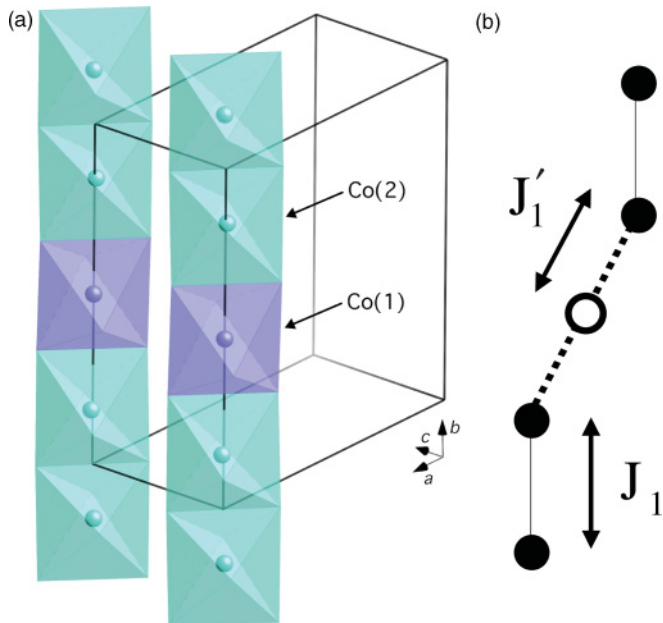


FIG. 1. (Color online) (a) Structure of triclinic CoV_2O_6 showing edge-sharing CoO_6 chains, with the two crystallographically independent sites indicated. For clarity, the V_2O_5 blocks are omitted. The principle exchange interactions in the chain are indicated in (b).

stirred until a gel formed. The gel was allowed to solidify then decomposed at 300°C for 3 h. The resulting solid was ground, pelleted, and heated at 600°C , 630°C , and 650°C for 12, 12, and 72 h, respectively. Powder x-ray diffraction using a Bruker D8-Advance showed a phase pure product. Magnetic susceptibility measurements were performed using a Quantum Design MPMS in varying fields up to 2 T as a function of temperature from 2 to 300 K in field- and zero-field-cooled conditions. Magnetization isotherms were recorded using the ac susceptibility option of a Quantum Design PPMS at temperatures from 2 to 35 K in fields of up to 9 T. This system was also used for specific heat measurements. For these, an additional measurement consisting of the sample mount and a small quantity of grease was performed before a small pellet of CoV_2O_6 was affixed. We used a variety of neutron powder diffractometers to characterize our samples. High-resolution data were collected at 2 K with wavelengths of 1.79 and 2.8 Å using the E9 instrument at HZB, Berlin.²⁵ A detector bank consisting of 64 ^3He detectors arranged at 2.5° intervals was employed. A total of 32 steps of the detector were carried out, resulting in a step size of 0.078° with a resolution minimum of $\Delta d/d \sim 2 \times 10^{-3}$ over a total angular range of 160° . The high-resolution data were refined by the Rietveld method using the GSAS suite of programs.²⁶ We also performed temperature-dependent measurements using the medium-resolution focusing diffractometer E6, also at HZB.²⁷ This instrument has a vertically and horizontally bent pyrolytic graphite monochromator and two large position-sensitive detectors giving a high flux at a wavelength of 2.4 Å. For both diffraction measurements, the sample was held in a 6-mm vanadium can and temperature was controlled using a standard Orange helium flow cryostat. Time-of-flight inelastic neutron scattering data were collected with IN5 at the

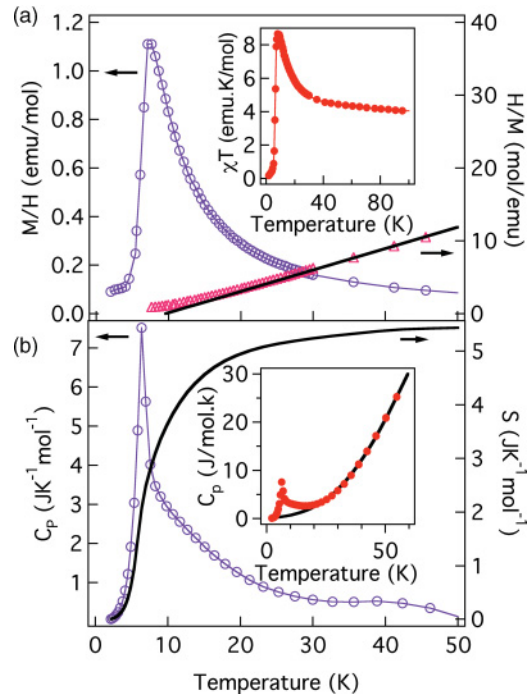


FIG. 2. (Color online) (a) Magnetic susceptibility and inverse susceptibility of CoV_2O_6 : the black line shows the Curie-Weiss fit to the high-temperature region described in text. Inset shows χT as a function of temperature. (b) Magnetic specific heat of CoV_2O_6 : line shows integrated magnetic entropy, inset shows total specific heat and estimated lattice contribution.

Institut Laue-Langevin, Grenoble. This instrument is a direct geometry cold neutron spectrometer with a 30-m^2 array of position-sensitive detectors. The sample was contained in an annular aluminum can placed in an Orange cryostat. All data were background corrected and normalized to the incoherent scattering of vanadium using standard routines in the program LAMP. We employed two configurations with incident energies of 3.27 and 7.08 meV, giving resolutions of around 0.1 and 0.35 meV, respectively.

III. RESULTS

A. Physical properties

The magnetic susceptibility of CoV_2O_6 , measured in a 500-Oe field, is shown in Fig. 2(a) and shows a sharp transition to an antiferromagnetically ordered state below $T_N = 7$ K. No divergence between field- and zero-field-cooled measurements was seen. The inverse susceptibility of CoV_2O_6 is well fitted by the Curie-Weiss law in the range 125–300 K with a fitted moment of $5.22(2)\mu_B$ and a Weiss temperature of $9.2(2)$ K. The moment shows a substantial increase due to spin-orbit coupling from the spin-only value for $S = 3/2$ ($3.87\mu_B$) as is often found for high-spin Co(II) in octahedral coordination with a $4T_{1g}$ ground state. The plot of χT versus T shown in the inset of Fig. 2(a) provides strong evidence for nearest-neighbor ferromagnetic correlations, as a sizable enhancement in χT is observed below 40 K, before an overall antiferromagnetic state is realized at T_N . These strong ferromagnetic correlations presumably originate in the CoO_6 chains, which have the shortest exchange pathway by far.

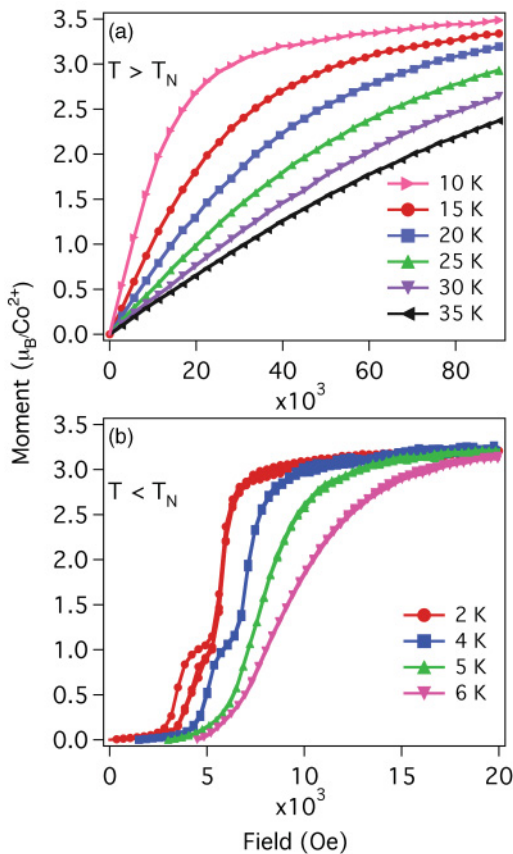


FIG. 3. (Color online) (a) Magnetization isotherms measured for CoV_2O_6 above magnetic ordering temperature; (b) magnetization isotherms measured for CoV_2O_6 below magnetic ordering temperature. A constant x -axis offset of 1500 Oe has been applied to the results in (b) for clarity. Lines in all plots are guides to the eye.

Long-range order is therefore the result of antiferromagnetic interchain (or plane) exchange.

The measured specific heat of a polycrystalline pellet between 2 and 70 K is shown in Fig. 2(b). A sharp λ anomaly is seen at 6.3 K, showing that only one magnetic transition occurs in this temperature range. Application of a 1-T field (which is above the metamagnetic field) both suppresses and broadens the peak (not shown). Differences between the specific heat in 0- and 1-T applied field were found up to 30 K, which suggests that magnetic fluctuations persist to at least this temperature, in agreement with the magnetic measurements reported above. We estimated the lattice contribution to the specific heat of CoV_2O_6 by fitting the high-temperature ($T > 30$ K) specific heat to a function of the form $AT^3 + BT^5$, the results of which are shown in Fig. 2(b) as a bold line. The magnetic entropy was obtained by subtracting the lattice contribution: the total obtained (5.43 J/mol K) is very close to the expected value for an $S = 1/2$ spin ($R \ln 2 = 5.76$ J/mol K). This result implies that the ground state of the Co^{2+} cations is well described by a Kramers doublet, and that strong single-ion anisotropy dominates the magnetic properties in the temperature region studied here. We note that 65% of the entropy is obtained above T_N , which confirms the low-dimensional nature of CoV_2O_6 .

$M(H)$ isotherms were recorded at a range of temperatures in the magnetically ordered region and above T_N in fields of up

to 9 T. Above T_N , the $M(H)$ isotherms were characteristic of a soft ferromagnet [Fig. 3(a)]. These observations corroborate the evidence from the susceptibility measurements for nearest-neighbor ferromagnetic exchange in CoV_2O_6 . Based on the isotherms shown in Fig. 3(a), these correlations can be seen to extend well above T_N , to ~ 35 K. Isotherms measured in the range $2 < T < T_N$ are shown in Fig. 3(b). At 2 K, two field-induced transitions are seen. The first transition at 3600 Oe corresponds to a plateau at almost exactly one third of the saturation magnetization ($0.95 \mu_B$), and the second transition at 5900 Oe is a metamagnetic transition to a plateau that shows the full saturation magnetization, $2.9 \mu_B$ at 9 T ($\mu_{\text{sat}} = 2.5 S = 3 \mu_B$). Considerable hysteresis was observed around the $1/3$ plateau at the lowest temperatures measured. $M(H)$ isotherms at higher temperatures show that the $1/3$ plateau becomes nonhysteretic at 4 K and disappears between 4.5 and 5 K [Fig. 3(b)], whereas the metamagnetic transition persists up to the Néel temperature.

B. Neutron powder diffraction measurements

Our neutron powder diffraction measurements at 2 K showed the presence of magnetic order and were also used to refine the crystallographic structure. The data were analyzed as follows. First, the crystal structure of CoV_2O_6 was refined from the short-wavelength (1.7973 Å) data collected on the high-resolution diffractometer E9. Due to the low symmetry, the lattice parameters and background were first refined using a Le Bail parameterless fit, giving results similar to those reported at room temperature. The peak shape parameters (a pseudo-Voigt function), scale function, and detector zero-point shift were then refined before the internal coordinates were allowed to vary. In order to reduce the number of variables, and because of the low scattering length of vanadium, the atomic displacement parameters (ADPs) of the Co and V sites were constrained to be equal, as were the oxygen ADPs. The possibility of antisite disorder between Co and V was investigated but failed to give an improvement in the fit residuals and was therefore discounted. The refinement for this histogram converged with $R_{wp} = 0.0664$ and $R_p = 0.0512$ and gave the results shown in Tables I and II. Notably, the refined Co-O-Co bond angles support our suggestion of ferromagnetic superexchange interactions along the chain direction, as in all cases, these are less than $\sim 95^\circ$. The observed, calculated, and difference plots for the Rietveld fits are shown in Figs. 4(a) and 4(b).

At low angles, magnetic reflections which did not index on the chemical unit cell were observed [Fig. 4(a)]. Previous work on related Mn- and CuV_2O_6 materials^{24,28} has shown that simple collinear ground states consisting of ferromagnetic chains coupled antiferromagnetically are found. However, neither of these compounds has strong single-ion anisotropy, which is expected to complicate the magnetic ordering. In the case of CoV_2O_6 , we were able to index the largest magnetic reflections on a supercell with $k = (1/2, 0, 0)$. A simple magnetic structure of ferromagnetic planes coupled antiferromagnetically accounted for the majority of the magnetic scattering. However, we also detected a large number of extra satellite reflections [see inset of Fig. 4(a)]. These resisted all attempts at indexing due to the low crystallographic symmetry

TABLE I. Refined atomic coordinates for CoV_2O_6 at 2 K from neutron powder diffraction. Lattice parameters were $a = 7.172(1)$, $b = 8.875(1)$, $c = 4.8088(8)$ Å, $\alpha = 90.288(2)$, $\beta = 93.931(2)$, $\gamma = 102.160(1)^\circ$. Refined ADPs are $\text{Co/V} = 0.2(1)$, $\text{O} = 0.73(3) \times 10^2$ Å².

Atom	x	y	z
Co(1)	0	0.5	0
Co(2)	0.0284(29)	0.1715(25)	0.028(4)
V(1)	0.704(16)	0.951(16)	0.462(24)
V(2)	0.719(17)	0.631(15)	0.433(24)
V(3)	0.498(16)	0.240(14)	0.143(21)
O(1)	0.1620(14)	0.4970(11)	0.3456(19)
O(2)	0.8413(15)	0.6399(13)	0.1623(21)
O(3)	0.1746(13)	0.6964(13)	0.8841(19)
O(4)	0.1537(13)	0.0250(12)	0.8270(19)
O(5)	0.1697(14)	0.8925(11)	0.3428(21)
O(6)	0.7877(12)	0.8019(14)	0.6320(18)
O(7)	0.4748(14)	0.9122(12)	0.7127(20)
O(8)	0.4724(14)	0.5803(12)	0.7046(21)
O(9)	0.5222(13)	0.7539(14)	0.2123(18)

and probably correspond to a long-wavelength modulation of the magnetic structure as seen²⁹ in, e.g., $\text{Ca}_3\text{Co}_2\text{O}_6$. Incommensurate magnetic structures, often with multiple propagation vectors, are common in systems with competing interactions. In this case, interplay between superexchange and single-ion anisotropy on the two crystallographically independent sites is a possible origin. Future single-crystal diffraction studies are necessary to solve this structure.

In order to determine the extent of critical fluctuations in CoV_2O_6 , we measured the intensity of several fundamental magnetic reflections as a function of temperature using E6. We chose the reflections between 30° and 40° marked in Fig. 4(a). A great deal of diffuse scattering was observed around the most intense magnetic reflection ($\sim 11^\circ$) near T_N , precluding use of this region of the diffraction patterns. A plot of the resulting normalized magnetic intensity versus reduced temperature is shown in Fig. 5 (assuming a Néel temperature of 6.3 K). A crossover between critical behavior and true long-range order is clearly seen at ~ 5.1 K, which matches

TABLE II. Selected refined distances and angles for CoV_2O_6 at 1.6 K from neutron powder diffraction.

	Bond distance or angle (Å or deg)
Co(1)-O(1) \times 2	1.962(9)
Co(1)-O(2) \times 2	2.036(11)
Co(1)-O(3) \times 2	2.024(11)
Co(2)-O(2)	1.993(25)
Co(2)-O(3)	2.113(25)
Co(2)-O(4)	2.005(23)
Co(2)-O(4)	2.099(25)
Co(2)-O(5)	2.193(21)
Co(2)-O(6)	2.009(21)
Co(1)-O(2)-Co(2)	94.8(8)
Co(1)-O(3)-Co(2)	91.6(7)
Co(2)-O(4)-Co(2)	93.5(9)

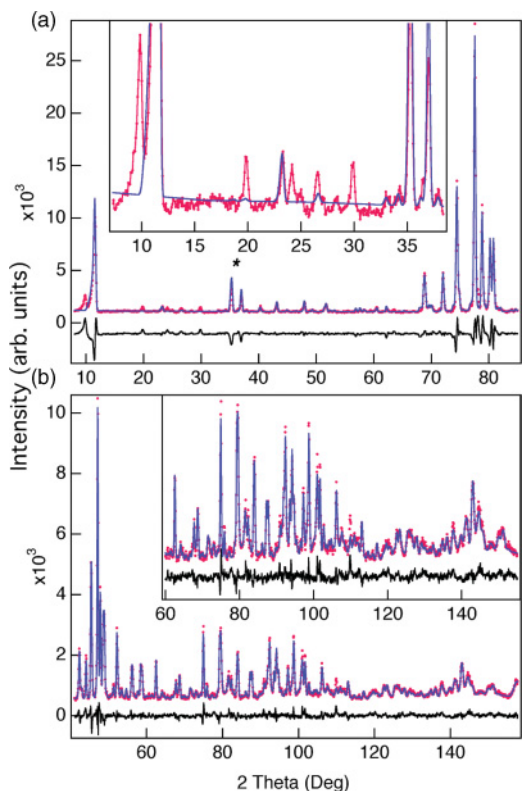


FIG. 4. (Color online) Observed, calculated, and difference plots for Rietveld fit to (a) 2.8 Å and (b) 1.79 Å neutron powder diffraction profiles of CoV_2O_6 at 2 K. Insets show unindexed magnetic satellite reflections and fit to high-angle data, respectively. The star in (a) marks the magnetic reflections followed to determine the order parameter using E6.

the region identified by the $M(H)$ isotherms without a $1/3$ magnetization plateau. A fit of a power-law function gave a critical exponent, $\beta = 0.22(2)$, which is at the upper end of the range found for anisotropic layered Ising systems,³⁰ although strongly reduced from the value³¹ of 0.35 found for MnV_2O_6 . Future measurements with better temperature resolution are needed to determine a more accurate value.

C. Inelastic neutron scattering measurements

A summary of the measurements performed with $E_i = 7.08$ meV and $T > T_N$ is shown in Fig. 6. Here the excitations are expected to be representative of the one-dimensional chains. As expected from the single-ion anisotropy, the majority of the inelastic spectral weight is gapped at 7 K. Due to the almost nondispersive nature of the magnetic response, the data are integrated over all momentum transfers and hence approximate to the magnetic density of states. The sharpness of the gapped peaks is significant, as Van Hove singularities are expected in the magnetic density of states in low-dimensional materials. This reinforces our observations from the specific heat measurements which showed that only 35% of the magnetic entropy is recovered at T_N , and we conclude that CoV_2O_6 is highly one-dimensional. Next, by following the temperature dependence of the inelastic response, we confirm the magnetic nature of all these features and show the origin of those above and below 1.5 meV is rather different.

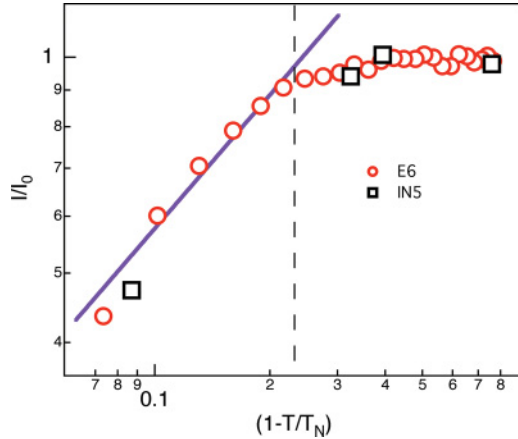


FIG. 5. (Color online) Temperature dependence of the elastic magnetic diffraction peaks recorded using E6. The black squares are elastic data recorded using IN5 with 5-Å incident neutrons. The solid line marks a power-law fit, and the dashed line marks the approximate extent of the critical region.

In the inset to Fig. 6 we show the integrated inelastic response above and below 1.5 meV. On warming, the intensity of the ~ 1 -meV features increases at the expense of the gapped

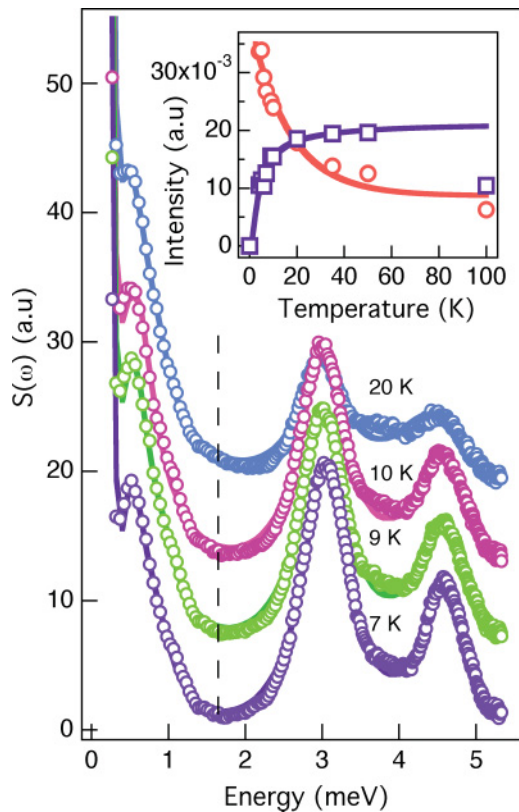


FIG. 6. (Color online) Inelastic neutron scattering from CoV_2O_6 for $7 < T < 20$ K. Data have been summed for all momentum transfers. The inset shows the temperature dependence of the integrated inelastic scattering above and below 1.5 meV. All lines are fits as described in the text.

bands up to at least 35 K, before collapsing. We fitted the data in this region to the gap function:

$$I \propto \exp(-E_g/k_B T). \quad (1)$$

The fitted gap value of 3.6(3) meV shows that this feature originates in neutrons scattering from thermally populated states in the gapped bands and that these are therefore soliton continuums rather than well-defined spin-wave excitations. The magnetic nature of the low-energy feature is confirmed by the plateau in intensity above 35 K. The drop in intensity at 50 K is caused by the redshift of the signal into the region below that used for the integration (i.e., the elastic line) and hence is not intrinsic. Structure also develops in the gapped bands on warming (principally a small increase in intensity at 3.8 meV). Future studies are needed to determine if this is also domain wall scattering or if it marks a crossover to a more spin-wave-like response as more crystal field levels are populated at higher temperatures. At temperatures far in excess of the magnetic exchange scale (200 K), the inelastic response disappears, to be replaced by quasielastic scattering showing that no phonons are present in this energy and momentum range (not shown).

We next examine in more detail the features at 7 K, including their energy [Fig. 7(a)] and Q dependencies [Fig. 7(b)]. The energies of the peaks shown in Fig. 7(a), extracted using the instrument resolution function and Lorentzian peak shapes, were 3.05(1) and 4.60(1) meV. For comparison, the asymmetric feature at low energy seen in Fig. 6 was peaked at 0.47(5) meV (fit not shown). In Fig. 7(b), the integral of the inelastic response for all energies [i.e., $S(Q)$] is shown. This approximately follows the squared Co^{2+} form factor (blue line) as expected for nearest-neighbor ferromagnetic (FM) exchange. The individual Q dependencies of the two features in $S(\omega)$ are in fact rather different [inset, Fig. 7(b)]. This helps to show that these are spin rather than crystal field excitations, which would follow the form factor. Crystal field excitations are also generally found at much higher energies in Co^{2+} -containing oxides.³² Taking into consideration the peculiar dimer arrangement of $\text{Co}(2)$ sites in CoV_2O_6 [Fig. 1(a)], and the peaked nature of the 4.60(1) meV feature, we performed an additional test against nearest-neighbor antiferromagnetic (AFM) coupling. We used the following powder averaged form factor:

$$S(Q) \propto f(Q)^2 [1 - \sin(Qd)/Qd]. \quad (2)$$

Here $f(Q)$ is the magnetic form factor and d is a characteristic distance. We also included a background term which accounts for the slow increase in intensity with Q found throughout our data. This is probably due to multiple incoherent scattering of phonons. The fitted value of d was 3.36(2) Å, which far exceeds the real value in the chains (~ 2.85 Å). This shows that CoV_2O_6 is not a loosely linked chain of dimer units and that the origins of the two features lie elsewhere.

To better define the thermally populated inelastic scattering below 1.5 meV, we show the results of the measurements in the low-energy configuration of IN5 over a wide range in temperature in Fig. 8. To remove the intrinsic effects of

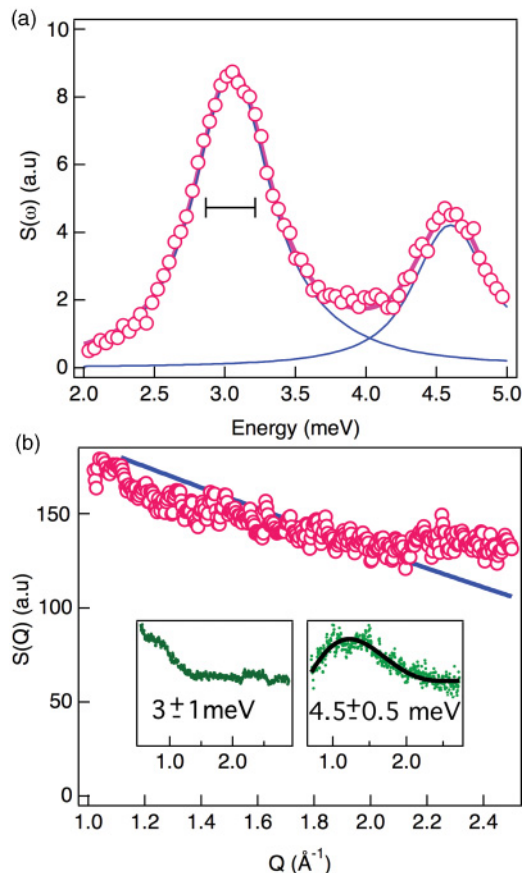


FIG. 7. (Color online) (a) Inelastic neutron scattering from CoV_2O_6 just above T_N at 7 K. Data have been summed for all momentum transfers, the lines are fits as discussed in the text, and the horizontal bar shows the instrumental resolution. (b) Energy-integrated inelastic neutron scattering data for CoV_2O_6 . The solid line shows the squared Co^{2+} form factor. The insets show the Q dependencies of the two principle excitations.

temperature, we have employed the fluctuation dissipation theorem to extract the generalized susceptibility $\chi''(Q, \omega)$:

$$S(Q, \omega) \propto \frac{\chi''(Q, \omega)}{1 - \exp(-\hbar\omega/k_B T)}. \quad (3)$$

The low-energy signal is clearly resolved into two peaks just above T_N as shown by the cuts at $Q = 1.3 \pm 0.1 \text{ \AA}^{-1}$ in Fig. 8.

This response reaches a maximum at ~ 11 K and collapses with further heating. In this region the data were fitted by the sum of two Lorentzians and a sloping background. The position of the excitations was $0.55(1)$ and $1.06(1)$ meV at 11 K. Upon heating, a gradual broadening and convergence of the two peaks was observed. Above 20 K, the data could be equivalently described by a single damped peak. These low-energy features thus approximately match the temperature region where our magnetic susceptibility and specific heat measurements detect low-dimensional ferromagnetic fluctuations (Fig. 2).

The onset of magnetic order, and the associated internal field in CoV_2O_6 , dramatically changes both the low- and high-energy spin excitations. As shown in Fig. 8, the former almost entirely disappears below 5 K. The gapped features are

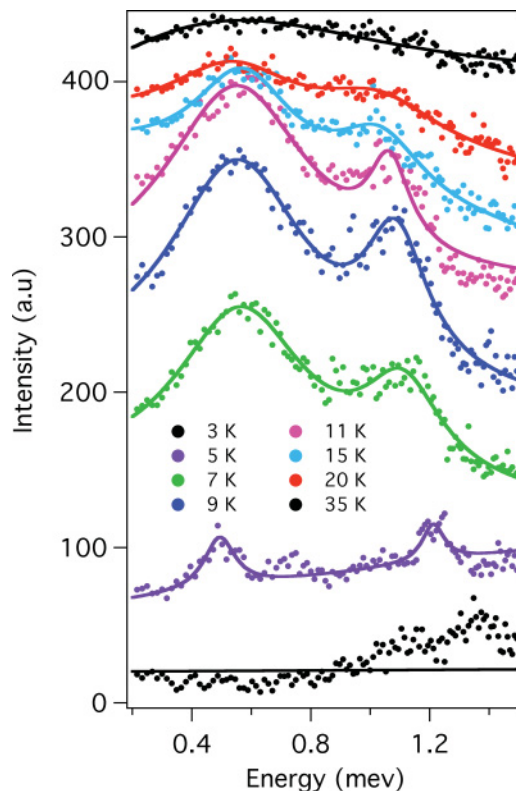


FIG. 8. (Color online) The results of the inelastic neutron scattering experiment using the low-energy configuration of IN5 for a range of temperatures. The data are presented in the form of constant $Q = 1.3 \pm 0.1 \text{ \AA}^{-1}$ and have been corrected for the effects of temperature using Eq. (3). The lines show the results of fitting Lorentzian functions and a sloping background.

also strongly affected, as shown by the data collected in the higher-energy configuration. Figure 9 shows that, below T_N , a more complex structure emerges from the broad bands at 3 and 4.5 meV. The two bands of excitations found above T_N split into at least five distinct modes, whose fitted positions (extracted using Lorentzian functions) are shown in the inset to Fig. 9. Assuming these are bound states, as discussed more fully below, an approximately linear increase in energy with the bound-state level was found (inset, Fig. 9).

IV. DISCUSSION

The above results represent a comprehensive investigation of the structure, magnetic properties, and dynamics of CoV_2O_6 . Our principal conclusion is that CoV_2O_6 may be regarded as a one-dimensional spin chain with considerable Ising-type anisotropy and ferromagnetic nearest-neighbor exchange. This is supported by both physical property and neutron scattering measurements. Here, we briefly review the former before commenting in detail on the latter and the phase diagram which we derive for CoV_2O_6 . Turning first to the physical property measurements in low magnetic fields, both the magnetic susceptibility [Fig. 2(a)] and the specific heat of CoV_2O_6 [Fig. 2(b)] show a single antiferromagnetic transition at $T_N = 6.3$ K. However, the positive Weiss temperature of $9.2(2)$ K and the peak seen in χT versus T above T_N imply nearest-neighbor ferromagnetic exchange. Our specific heat

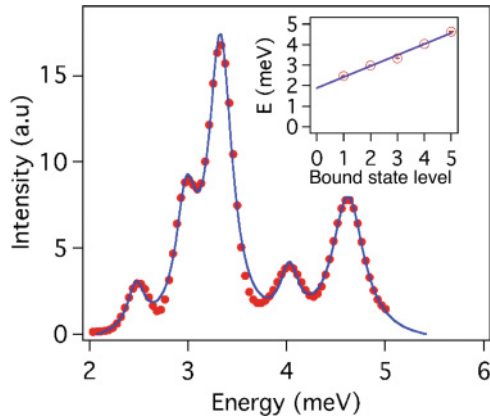


FIG. 9. (Color online) Momentum transfer summed inelastic scattering of CoV_2O_6 at 1.5 K and 7.08 meV incident energy showing the confining effect of the internal field in the magnetically ordered state. Lines are fits to Lorentzian functions. The inset shows the energy of the bound states with a linear fit.

measurements confirm the presence of magnetic fluctuations up to temperatures of around 40 K. These measurements also provide the first piece of evidence for Ising anisotropy as the integrated magnetic entropy is very similar to that expected for an effective $S = 1/2$ spin. This picture is strongly confirmed by the $M(H)$ isotherms shown in Figs. 3(a) and 3(b) and the critical exponent $\beta = 0.22(2)$ derived from neutron powder diffraction (Fig. 5). These physical property measurements compare well with those previously reported^{33,34} for Ising FM salts such as $\text{CoCl}_2 \cdot 2\text{H}_2\text{O}$, which also show a $1/3$ magnetization plateau.

Our results start to differ from those reported for these simple materials as we consider our neutron scattering data. First, our diffraction measurements at 2 K show that the magnetic order is incommensurate, although the majority of the scattering can be modeled assuming FM chains coupling antiferromagnetically (Fig. 4). Second, as we now discuss in more detail, our inelastic neutron scattering results do not coincide with predictions for a linear FM Ising spin chain. Unlike AFM Ising chains, soliton excitations in materials with a FM nearest-neighbor exchange J_1 do not deconfine. The excitations are instead isolated flips of blocks of spins,⁶ and the results of zero-field inelastic neutron scattering may be modeled using standard spin-wave theory.³⁵ The degeneracy of these excitations may also be lifted by applying magnetic fields, and the resulting spectrum of spin flips has also been observed using infrared transmission experiments.³⁶ The observation of sharp, almost dispersionless features in CoV_2O_6 , as well as temperature-induced scattering, is inconsistent with these expectations. This shows that any Hamiltonian proposed for CoV_2O_6 must include terms which account for the low crystallographic symmetry. Of obvious importance here is the period 3 variation of crystallographically distinct Co sites (Fig. 1). This may result in a variation in the local easy axis on moving along the chain. In a simple model based on band folding, the presence of two sites would also result in two gapped soliton bands as experimentally observed. Furthermore, this may also be consistent with the presence of two low-energy features. The fitted gap function for these

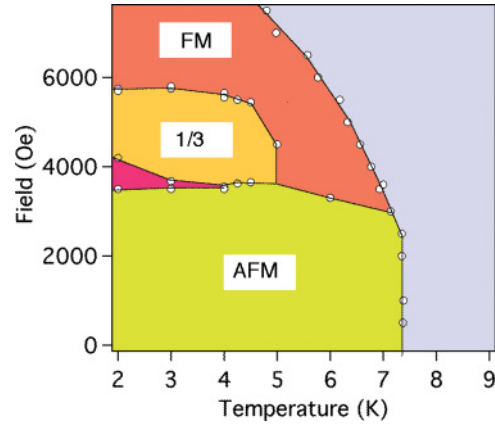


FIG. 10. (Color online) The temperature-magnetic field phase diagram of CoV_2O_6 established from the physical property measurements in this work. The various phases are labeled as FM (fully polarized ferromagnetic), $1/3$ (plateau in magnetization at $1/3$ of saturation), and AFM (antiferromagnetic phase). The region of hysteresis found at the lower bound of the $1/3$ plateau is shown in magenta (unlabeled dark shading).

shows that they originate in Villain-mode-type scattering in the gapped bands. As both show the same temperature dependence, the doubling may hence be an indication of both intra- and interband scattering.

On cooling below T_N , the complex features which emerge from the gapped bands between 2 and 5 meV are similar to what was reported for the related compound CoNb_2O_6 . This is proposed to be a result of the confining effect of the internal field caused by magnetic order on domain walls. Interchain coupling imposes a linearly increasing potential between propagating domain walls, quantizing the continuum into a ladder of bound states^{5,37} above the energy required to create an isolated spin flip ($2J_1$). The overall picture seen in CoV_2O_6 (inset, Fig. 9) qualitatively agrees with this picture; however, significant dispersion is not evident from our data. In Ref. 5, an empirical parameter which tuned the dispersion was introduced into the Hamiltonian. This is strongly dependent on the exchange topology in each individual material. In our powder averaged case, the energies of the bound states increase linearly, and from the y intercept we estimate an average intrachain interaction of $0.95(1)$ meV.

The phase diagram of CoV_2O_6 that we have determined from our physical property measurements is shown in Fig. 10. Of particular note is the $1/3$ magnetization plateau. This feature is not found for all temperatures within the antiferromagnetically ordered region below 6.3 K, instead setting in below ~ 5 K. Interestingly, the region without a plateau coincides extremely well with the region of critical fluctuations identified by neutron powder diffraction shown in Fig. 5. Examination of Fig. 8 also shows that a small amount of the thermally populated domain wall inelastic scattering also remains down to 5 K. This is in some respects similar to the situation found in the intermediate partially disordered magnetic structure in CsCoCl_3 . This compound has a coexistence of $2/3$ ordered spins and soliton excitations in a low-temperature region before complete order sets in.³⁸ A plausible explanation of our results for CoV_2O_6 , consistent with the anisotropic coupling between chains, would be a crossover between two-

and three-dimensional (2D and 3D) magnetic order below 5 K.

V. CONCLUSIONS

We have reported the bulk synthesis and characterization of powder samples of CoV_2O_6 . We have used a combination of physical property and neutron scattering measurements to show that CoV_2O_6 is a one-dimensional Ising magnet with ferromagnetic nearest-neighbor exchange. In the ordered state, we have discovered a crossover from a metamagnet with strong fluctuations between 5 K and T_N to a state with a 1/3 magnetization plateau at $2 < T < 5$ K. This may arise from a dimensionality crossover from 2D to 3D magnetic ordering. The temperature dependence of the inelastic neutron scattering signal shows that the principle excitations are domain walls and at low temperatures we detect signatures

of confinement caused by the internal field. It is likely that the low symmetry of CoV_2O_6 , especially the presence of two crystallographically independent sites in the edge-sharing chains, strongly influences the magnetic properties.

Significant challenges remain, including resolving the exact magnetic structure adopted below 7 K and a complete modeling of the inelastic neutron spectra. We anticipate that these can be addressed once single crystals of CoV_2O_6 become available.

ACKNOWLEDGMENTS

S.A.J.K. and J.P.A. thank the EPSRC and Leverhulme trust, respectively, for support. We additionally acknowledge the Helmholtz Zentrum Berlin for funding and the Institute Max von Laue-Paul Langevin for access to their instruments. We thank D.A. Tennant for useful discussions.

*kimber@esrf.fr

†j.p.attfield@ed.ac.uk

¹D. A. Tennant, T. G. Perring, R. A. Cowley, and S. E. Nagler, *Phys. Rev. Lett.* **70**, 4003 (1993).

²S. E. Nagler, W. J. L. Buyers, R. L. Armstrong, and B. Briat, *Phys. Rev. B* **28**, 3873 (1983).

³P. Pfeuty, *Ann. Phys.* **57**, 79 (1970).

⁴S. Sachdev, *Quantum Phase Transitions* (Cambridge University Press, Cambridge, UK, 1999).

⁵R. Coldea, D. A. Tennant, E. M. Wheeler, E. Wawrzynska, D. Prabhakaran, M. Telling, K. Habicht, P. Smeibidl, and K. Kiefer, *Science* **327**, 177 (2010).

⁶J. Villain, *Physica B+C* **79**, 1 (1975).

⁷W. Scharf, H. Weitzel, I. Yaeger, I. Maartense, and B. M. Wanklyn, *J. Magn. Magn. Mater.* **13**, 121 (1979).

⁸S. Mitsuda, K. Hosoya, T. Wada, H. Yoshizawa, T. Kanawa, M. Ishikawa, K. Miyatani, K. Saito, and K. Kohn, *J. Phys. Soc. Jpn.* **63**, 3568 (1994).

⁹I. Maartense, I. Yaeger, and B. M. Wanklyn, *Solid State Commun.* **21**, 93 (1977).

¹⁰T. Hanawa, K. Shinkawa, M. Ishikawa, K. Miyatani, K. Saito, and K. Kohn, *J. Phys. Soc. Jpn.* **63**, 2706 (1994).

¹¹C. Heid, H. Weitzel, P. Burllet, M. Bonnet, W. Gonschorek, T. Vogt, J. Norwig, and H. Fuess, *J. Magn. Magn. Mater.* **151**, 123 (1995).

¹²C. Heid, H. Weitzel, P. Burllet, M. Winkelmann, H. Ehrenberg, and H. Fuess, *Physica B* **234**, 547 (1997).

¹³S. Kobayashi, S. Mitsuda, M. Ishikawa, K. Miyatani, and K. Kohn, *Phys. Rev. B* **60**, 3331 (1999).

¹⁴S. Kobayashi, S. Mitsuda, T. Jogetsu, J. Miyamoto, H. Katagiri, and K. Kohn, *Phys. Rev. B* **60**, 9908 (1999).

¹⁵S. Kobayashi, S. Mitsuda, and K. Prokes, *Phys. Rev. B* **63**, 024415 (2000).

¹⁶H. Weitzel, H. Ehrenberg, C. Heid, H. Fuess, and P. Burllet, *Phys. Rev. B* **62**, 12146 (2000).

¹⁷H. Fjellvag, E. Gulbrandsen, S. Aasland, A. Olsen, and B. C. Hauback, *J. Solid State Chem.* **124**, 190 (1996).

¹⁸H. Kageyama, K. Yoshimura, K. Kosuge, H. Mitamura, and T. Goto, *J. Phys. Soc. Jpn.* **66**, 1607 (1997).

¹⁹A. Maignan, V. Hardy, S. Hebert, M. Drillon, M. R. Lees, O. Petrenko, D. M. Paul, and D. Khomskii, *J. Mater. Chem.* **14**, 1231 (2004).

²⁰B. Jasper-Tönnies and Hk. Müller-Buschbaum, *Z. Anorg. Allg. Chem.* **508**, 7 (1984).

²¹H.-K. Müller-Buschbaum and M. Kobel, *Z. Anorg. Allg. Chem.* **596**, 23 (1991).

²²Z. He, J.-I. Yamaura, Y. Ueda, and W. Cheng, *J. Am. Chem. Soc.* **131**, 7554 (2009).

²³A. N. Vasilev, L. A. Ponomarenko, A. I. Smirnov, E. V. Antipov, Y. A. Velikodny, M. Isobe, and Y. Ueda, *Phys. Rev. B* **60**, 3021 (1999).

²⁴S. A. J. Kimber and J. P. Attfield, *Phys. Rev. B* **75**, 064406 (2007).

²⁵D. M. Többens, N. Stüßer, K. Knorr, H. M. Mayer, and G. Lampert, *Mater. Sci. Forum* **378-381**, 288 (2001).

²⁶A. Larson and R. V. Dreele, Los Alamos National Laboratory Report No. LAUR 86-748, 1994 (unpublished).

²⁷N. Stüßer and M. Hofmann, *Nucl. Methods A* **482**, 744 (2002).

²⁸J. Kikuchi, I. Ishiguchi, M. Motoya, I. Itoh, I. Inari, E. Eguchi, and A. Akimitsu, *J. Phys. Soc. Jpn.* **69**, 2660 (2000).

²⁹S. Agrestini, L. C. Chapon, A. Daoud-Aladine, J. Schefer, A. Gukasov, C. Mazzoli, M. R. Lees, and O. A. Petrenko, *Phys. Rev. Lett.* **101**, 097207 (2008).

³⁰A. Taroni, S. T. Bramwell, and P. C. W. Holdsworth, *J. Phys. Condens. Matter* **20**, 275233 (2008).

³¹S. A. J. Kimber, Ph.D. thesis, University of Edinburgh, 2007.

³²T. Nguyen, S. E. Nagler, R. A. Cowley, T. Perring, and R. Osborn, *J. Phys. Condens. Matter* **7**, 2917 (1995).

³³W. Schneider and H. Weitzel, *Solid State Commun.* **13**, 303 (1973).

³⁴H. Weitzel and W. Schneider, *Solid State Commun.* **14**, 1025 (1974).

³⁵W. Montfrooij, G. E. Granroth, D. G. Mandrus, and S. E. Nagler, *Phys. Rev. B* **64**, 134426 (2001).

³⁶J. B. Torrance and M. Tinkham, *Phys. Rev.* **187**, 595 (1969).

³⁷J. P. Goff, D. A. Tennant, and S. E. Nagler, *Phys. Rev. B* **52**, 15992 (1995).

³⁸J. P. Boucher, L. P. Regnault, J. Rossat-Mignod, Y. Henry, J. Bouillot, and W. G. Stirling, *Phys. Rev. B* **31**, 3015 (1985).