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Magnons and continua in a magnetized and dimerized spin-1/2 chain

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We examine the magnetic field dependent excitations of the dimerized spin-1/2 chain, copper nitrate, with antiferromagnetic intra-dimer exchange $J_1 = 0.44(1)$ meV and exchange alternation $\alpha = J_2/J_1 = 0.26(2)$. Magnetic excitations in three distinct regimes of magnetization are probed through inelastic neutron scattering at low temperatures. At low and high fields there are three and two long-lived magnon-like modes, respectively. The number of modes and the anti-phase relationship between the wave-vector dependent energy and intensity of magnon scattering reflect the distinct ground states: A singlet ground state at low fields $\mu_0 H < \mu_0 H_{c1} = 2.8$ T and an $S_z = 1/2$ product state at high fields $\mu_0 H > \mu_0 H_{c2} = 4.2$ T. In the intermediate field regime, a continuum of scattering for $\hbar\omega \approx J_1$ is indicative of a strongly correlated gapless quantum state without coherent magnons.

I. INTRODUCTION

Insulating solids containing dimers of magnetic atoms with antiferromagnetic interactions can form a non-magnetic singlet ground state. As opposed to conventional dielectrics, the incipient magnetism of inter-atomic singlets can be exposed by chemical doping, pressure, and magnetic fields and is accessible to inelastic thermal neutron scattering. There are thus excellent opportunities to explore quantum collective phenomena at the onset of magnetism in such materials.

A case in point is materials containing dimerized antiferromagnetic spin-1/2 chains.^{1–5} When antiferromagnetic Heisenberg interactions prevail ($J_1 \gg J_2 \equiv \alpha J_1$), these spin systems can be driven through *two* quantum critical points, both associated with Bose-Einstein condensation of magnons.^{6–15} Immediately above the lower critical field, $\mu_0 H_{c1} \approx J_1(1 - \alpha)/(g\mu_B)$, a macroscopic density of magnons gives rise to magnetization.^{16–19} The reverse situation pertains at the upper critical field where magnon condensation *below* $\mu_0 H_{c2} = J_1(1 + \alpha)/(g\mu_B)$ reduces the magnetic moment per spin below saturation ($M < M_{\text{sat}} \equiv g\mu_B/2$).

While the intermediate field state for the idealized alternating spin chain is predicted to be a Luttinger liquid with dynamic transverse staggered spin correlations,^{20–22} the quantum critical nature of this phase implies sensitivity to sub-leading interactions and anisotropies. These ultimately control the low temperature behavior of real materials for $H_{c1} < H < H_{c2}$ where Néel^{23,24} or spin-Peierls order²⁵ is expected at sufficiently low temperatures.

A number of spin-1/2 dimer compounds have been subjected to detailed thermo-magnetic measurements exploring the full field range.^{15,26–34} To understand spin correlations at the atomic scale, however, requires inelastic neutron scattering experiments, which so far have been limited to the vicinity of the lower critical field H_{c1} .^{35–43} By selecting a material, copper nitrate, with

a suitable energy scale and utilizing high efficiency cold-neutron instrumentation, we can provide a comprehensive set of data for the field and wave vector dependence of collective magnetic excitations in a system of interacting spin-1/2 dimers throughout the relevant field range. Copper nitrate is an excellent experimental realization of an alternating chain 1d antiferromagnet. This is a specific example of a dimer system consisting of exchange coupled spins with an alternation in the magnitude of exchange interactions between nearest neighbor spin sites. Although there are other experimental examples of antiferromagnetic alternating chains including $(\text{VO})_2\text{P}_2\text{O}_7$,^{44–47} CuWO_4 ,⁴⁸ and $\text{Cu}(\text{L-aspartato})(\text{H}_2\text{O})_2$,⁴⁹ the energy scale of the triplet excitation in copper nitrate makes it well suited for studies as a function of applied magnetic fields.

While the resolution of the present experiment is insufficient to separate a presumed low energy continuum from intense elastic nuclear scattering, the experiment provides evidence for an extended gapless phase for $H_{c1} < H < H_{c2}$ in the form of a continuum of scattering for $J_1(1 - \alpha) < \hbar\omega < J_1(1 + \alpha)$. This is direct experimental evidence for a critical spin liquid state in the intermediate field regime. The paper is organized as follows. Section II provides background information about copper nitrate including new results on inter-chain interactions. Section III describes sample preparation and neutron instrumentation. The experimental data from complementary triple axis and time of flight neutron spectrometers are presented in section IV. Focusing in succession on each regime of the phase diagram, section V discusses the data. An exact diagonalization study, which is a useful point of reference opens this section. A summary of the conclusions from this work is presented in section VI.

II. COPPER NITRATE

A. Alternating spin chain model

The strongest interactions linking spin-dimers in copper nitrate ($\text{Cu}(\text{NO}_3)_2 \cdot 2.5\text{D}_2\text{O}$, (CN)) create a quasi-one-dimensional spin system that can be described as an alternating 1d antiferromagnetic chain. In an external magnetic field $\mathbf{H} = H\hat{z}$, the spin-Hamiltonian is

$$\mathcal{H} = \sum_n J_1(\mathbf{S}_{2n} \cdot \mathbf{S}_{2n+1} + \alpha \mathbf{S}_{2n+1} \cdot \mathbf{S}_{2n+2}) - g\mu_B\mu_0 \mathbf{H} \cdot \mathbf{S}_n, \quad (1)$$

where J_1 is the intra-dimer exchange, α is the ratio of inter- and intra-dimer exchange $\alpha = J_2/J_1$, g is the Landé g-factor, μ_B is the Bohr magneton, and μ_0 is the vacuum permeability.

Early thermo-magnetic measurements on CN showed exponentially activated ($\exp(-\Delta/T)$) susceptibility and specific heat data indicative of an energy gap, Δ , in the magnetic excitation spectrum.^{50–52} These results and nuclear magnetic resonance (NMR) measurements⁵³ were analyzed in terms of isolated dimers with weak inter-dimer interactions. The first accounts of inter-dimer interactions creating a 1d system resulted from an analysis of the crystal structure,⁵⁴ and additional thermo-magnetic studies.⁵⁵ Two models emerged: an alternating chain model and a two-leg ladder model. The alternating chains were proposed to extend along the [101] direction, while the crystalline b -axis was proposed as the extended direction for the 2-leg ladder model. The originally proposed alternating chain was mistakenly projected onto the [101] axis. Both models identify the spin-dimer as composed of nearest neighbor copper sites with the intra-dimer vector \mathbf{d}_1 (Fig. 1(c)). NMR measurements as a function of applied field and orientation showed the inter-dimer exchange primarily lies in the ac plane forming alternating chains rather than spin-ladders extending along b .⁵⁶

Specific heat and NMR studies of CN provide evidence of a field induced phase transition to long range antiferromagnetic order for $T \leq 175$ mK in magnetic fields between $\mu_0 H_{c1} = 2.8$ T and $\mu_0 H_{c2} = 4.2$ T.⁵⁷ A model to account for these phase transitions was developed by mapping the system in the intermediate field regime onto a $S = \frac{1}{2}$ linear chain in an effective magnetic field H_E , such that $H_E = 0$ when $H = \frac{1}{2}(H_{c1} + H_{c2})$.⁵⁸ Subsequent thermo-magnetic measurements were interpreted with moderate success based on this approach.^{56,59} An elastic neutron scattering experiment investigating the ordered phase of CN supported the alternating chain model and confirmed the ordered magnetic structure proposed by Diederix *et al.*⁶⁰ However, as we shall see in section II B, Diederix's alternating chain model incorrectly placed the chain axis along the [101] direction.

The aforementioned measurements led to general acceptance of the alternating chain model for CN, and established the exchange parameters to be given by $J_1 \approx$

0.44 meV and $\alpha \approx 0.24$. A direct spectroscopic measurement of the magnetic excitation spectrum was subsequently reported by Xu *et al.*⁶¹ These measurements showed the dynamic spin correlation function $\mathcal{S}(\tilde{q}, \hbar\omega)$ of the singlet-triplet excitations is well described by the single mode approximation (SMA) with dispersion relation

$$\epsilon(\mathbf{Q}) = J_1 - \sum_{\mathbf{u}} \frac{J_{\mathbf{u}}}{2} \cos(\mathbf{Q} \cdot \mathbf{u}). \quad (2)$$

Here J_1 is the intra-dimer exchange and $J_{\mathbf{u}}$ the inter-dimer exchange.⁶¹ Xu *et al.* further determined that along with broadening of the excitations the bandwidth decreases as temperature is increased above $T \approx 1$ K. The band narrowing does not arise from changes in the actual exchange constants, but from magnon-magnon interactions and can be accounted for by the random phase approximation.³⁷ In this paper, we shall at times list effective exchange constants determined by fitting dispersion relations measured at temperatures above 1 K to Eq. 2 and its generalizations. Such parameters are related to the actual exchange constants through $\tilde{J}_2 = n(T)J_2$, where $n(T)$ is the singlet-triplet population difference as discussed by Xu *et al.* Such low T renormalization does not occur for J_1 .⁶¹ Wave-vector dependent two-magnon excitations were also observed at higher energy transfers using inelastic neutron scattering.⁶² More recently, the zero-field single and multimagnon interactions have been examined with high resolution inelastic neutron scattering techniques to illustrate the influence of temperature on the quasiparticle excitations.⁶³

B. Crystal Structure

CN is monoclinic with space group $I2/c$ ⁶⁴ and room temperature lattice constants $a = 16.45$ Å, $b = 4.94$ Å, $c = 15.96$ Å, and $\beta = 93.77^\circ$.^{54,65,66} We find the low-temperature, $T \leq 4$ K, lattice constants to be $a = 16.12$ Å, $b = 4.90$ Å, $c = 15.84$ Å, and $\beta = 92.90^\circ$. Depicted in Fig. 1, the crystal structure features Cu^{2+} ions in a distorted 4+1 tetrahedral coordination with five oxygen atoms, (Fig. 1(c)). The four basal plane Cu-O bonds each have a bond length of approximately 2 Å while the apical Cu-O bond length is 2.39 Å.

The J_1 -coupled dimer is formed by neighboring Cu^{2+} ions where adjacent pairs of the short bond coordinated oxygen atoms form almost parallel planes that establish two Cu-O-H-O-Cu linkages as shown in Fig. 1(c). This exchange path is hydrogen bond mediated via a water group coordinated to one copper atom and a nitrate group coordinated to the other copper atom.

The alternating chain model involves structures extending along the [111] and $[\bar{1}\bar{1}\bar{1}]$ directions with Cu-O-N-O-H-O-Cu interdimer contacts. As shown in Fig. 1(b) and schematically in Fig. 1(d)-(e), each $[10\bar{1}]$ plane contains interpenetrating [111] and $[\bar{1}\bar{1}\bar{1}]$ alternating chains. Though the sequence of atomic contacts are identical

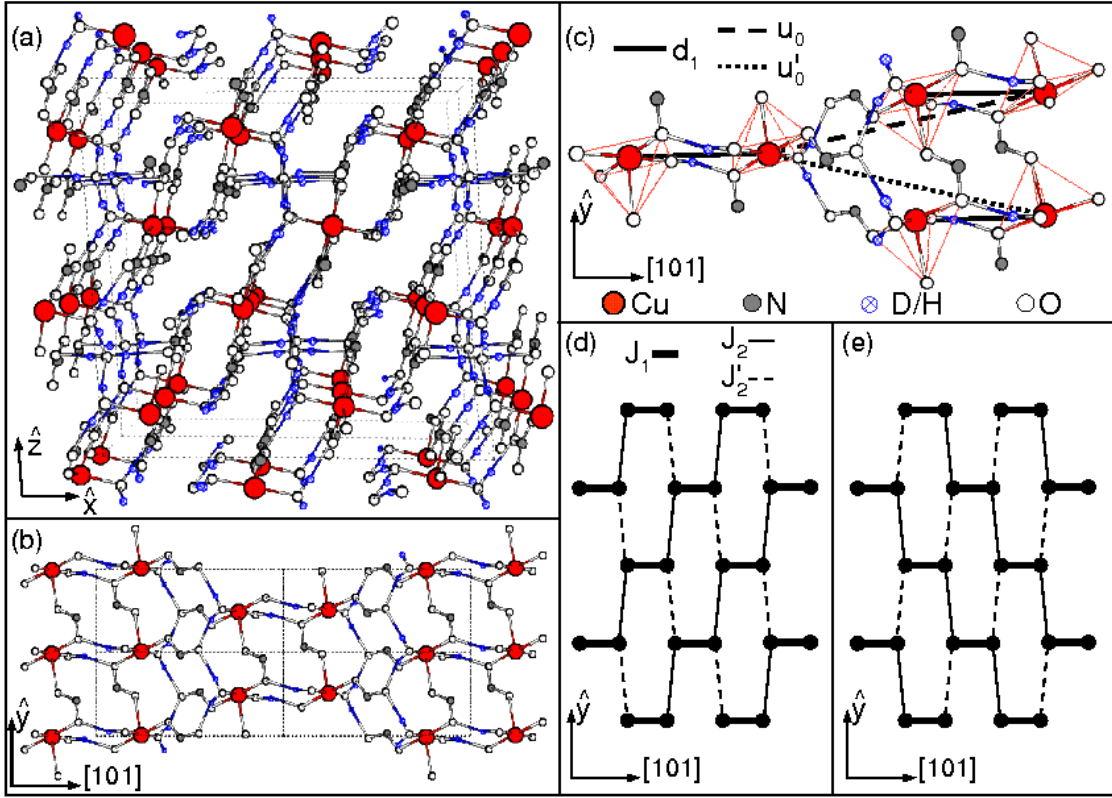


FIG. 1: (color online) Crystal structure of CN. (a) Perspective view along the b -axis. (b) View along the $[10\bar{1}]$ direction. (c) Expanded perspective view along the $[10\bar{1}]$ direction with 4+1 coordination of Cu^{2+} sites highlighted by square pyramidal polyhedra. Vectors \mathbf{d}_1 , \mathbf{u}_0 and \mathbf{u}'_0 are the respective intra- and two possible inter-dimer vectors described in the text. The interdimer exchange path we propose connects dimer units for which the basal planes of the 4+1 coordination are nearly co-planar along the $[111]$ direction in (b), (c) and (d) via the \mathbf{u}_0 interdimer bond and illustrated by J_1 - J_2 alternating chains in (d) and (e). (d) and (e) Schematic of interdimer interactions in CN in adjacent $[10\bar{1}]$ planes. Intra-dimer interactions, J_1 , are represented by circles linked by solid heavy lines. The two possible interdimer interactions are represented as solid (J_2) and dashed (J'_2) lines. In (b) and (c), proton sites D/H(4) and D/H(5), and oxygen sites O(3), O(4), and O(9) as enumerated in Ref. 54 have been omitted for clarity. These omitted atoms participate in bonding with water sites adjoining the copper atom, secondary exchange paths along the b direction, and bonding between the $[10\bar{1}]$ planes.

for both chains in a single $[10\bar{1}]$ plane, the bond angles and distances are different. We denote the two potential inter-dimer exchange constants as J_2 and J'_2 . Identical interactions for the two interpenetrating chains ($J_2 = J'_2$) would produce a two-dimensional distorted square lattice rather than alternating chains. In the alternating chain model, $J_2 \gg J'_2$, such that only one type of magnetic chain, $[111]$ or $[\bar{1}\bar{1}1]$, populates a single $[10\bar{1}]$ plane, while the other type of chain populates neighboring planes as shown in Figs. 1(d)-(e).

The fractional coordinates of the intra-dimer vector is $\mathbf{d}_1 = [0.25, \pm 0.03, 0.23]$ ($|\mathbf{d}_1| = 5.33 \text{ \AA}$), and the center to center inter-dimer vectors are \mathbf{u}_0 , $\mathbf{u}'_0 = [1, \pm 1, 1]/2$ ($|\mathbf{u}_0| = |\mathbf{u}'_0| = 11.351 \text{ \AA}$). The $[111]$ alternating chain in Fig. 1(c) (dashed line) connects dimer units with neighboring 4+1 oxygen coordinated basal planes that are nearly co-planar while the $[\bar{1}\bar{1}1]$ alternating chain in Fig. 1(c) (dotted line) connects dimer units with neighboring 4+1 oxygen coordinated basal planes that are nearly parallel but *not* co-planar.

Based on experience with other Cu^{2+} systems with similar coordination, and the fact that the dominant $3d$ electron density resides in the basal plane for this coordination,^{68–70} we hypothesize that the dashed exchange path, \mathbf{u}_0 , in Fig. 1(c), mediates the dominant inter-dimer exchange interaction. This corresponds to a nearest inter-dimer spin-spin distance of 6.22 \AA , as opposed to 6.32 \AA for the other exchange path. However, because the neighboring $[10\bar{1}]$ planes contain identically coordinated alternating chains in the other direction, neutron scattering can not distinguish between these two types of coordination. As discussed below, we are however, able to place limits on the amount of two-dimensional interchain exchange, J'_2 .

III. EXPERIMENTAL TECHNIQUES

A. Sample preparation

Deuterated CN single crystals were grown by cooling a saturated solution of deuterated CN in D_2O .⁶⁷ Deuterated CN was obtained through repeated distillation (3 - 4 times) of commercially available hydrogenous CN in high-purity D_2O ($> 99.9\%$). Seed crystals were initially grown by rapid cooling of saturated solutions from 80°C to 30°C . Single crystals were subsequently grown from suspended seeds in filtered saturated solutions of CN by slowly cooling from 50°C to 30°C . The sample used for inelastic neutron scattering consisted of three 94% deuterated single crystals with a total mass of 9.25 grams co-aligned to within 1.25° . A photograph of the sample assembly is shown in the Supplemental Material.⁷¹

B. Inelastic Neutron Scattering

Field dependent measurements were made using the SPINS triple-axis and the DCS time-of-flight spectrometers at NIST. The b -axis was vertical so that \mathbf{Q} was in the $(h0l)$ plane with equal contributions to the magnetic scattering from alternating chains extending along the $[111]$ and $[\bar{1}\bar{1}\bar{1}]$ directions. The sample environment for both measurements consisted of a dilution refrigerator within a split-coil 11.5 Tesla vertical field cryomagnet. Scattering intensities were normalized to the incoherent elastic scattering from a standard vanadium sample to provide a measurement in absolute units of the following two-point dynamic spin correlation function:

$$S^{\alpha\beta}(\mathbf{Q}, \omega) = \frac{1}{2\pi\hbar} \int_{-\infty}^{\infty} dt e^{-i\omega t} \frac{1}{N} \sum_{\mathbf{r}\mathbf{r}'} e^{i\mathbf{Q}\cdot(\mathbf{r}-\mathbf{r}')} \langle S_{\mathbf{r}}^{\alpha}(t) S_{\mathbf{r}'}^{\beta}(0) \rangle \quad (3)$$

Here $\hbar\omega$ and $\hbar\mathbf{Q}$ are energy and momentum transfer to the sample, α and β indicate cartesian components, N is the number of Cu^{2+} sites in the sample, and $S_{\mathbf{r}}^{\alpha}$ indicates the α component of the spin operator at position \mathbf{r} . In the following we shall use $\tilde{q} = \mathbf{Q} \cdot \mathbf{u}_0$ to indicate wave-vector transfer along the chains.

The normalized magnetic portion $\tilde{I}(\mathbf{Q}, \hbar\omega)$ of the inelastic neutron scattering intensity is related to the dynamic spin correlation function as follows

$$\tilde{I}(\mathbf{Q}, \hbar\omega) = \int d^3\mathbf{Q}' \hbar d\omega' \mathcal{R}_{\mathbf{Q},\omega}(\mathbf{Q} - \mathbf{Q}', \omega - \omega') \left| \frac{g}{2} F(\mathbf{Q}') \right|^2 \sum_{\alpha\beta} (\delta_{\alpha\beta} - \hat{Q}'_{\alpha} \hat{Q}'_{\beta}) S^{\alpha\beta}(\mathbf{Q}', \omega'), \quad (4)$$

where $F(\mathbf{Q})$ is the magnetic form factor of the Cu^{2+} ion and $\mathcal{R}_{\mathbf{Q},\omega}$ is the instrumental resolution function satisfying the following normalization condition.⁷³

$$\int d^3\mathbf{Q}' \hbar d\omega' \mathcal{R}_{\mathbf{Q},\omega}(\mathbf{Q} - \mathbf{Q}', \omega - \omega') = 1 \quad (5)$$

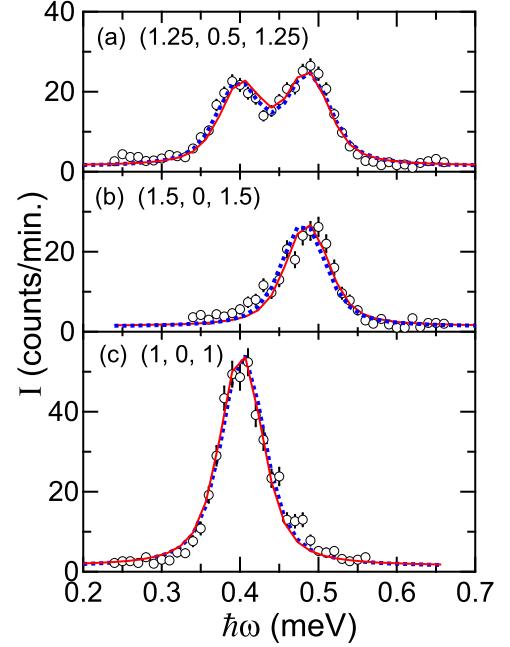


FIG. 2: (color online) Constant \mathbf{Q} scans for CN at $T = 1.25$ K. Red solid lines are from a global fit to resolution convolved single modes with dispersion as listed in Eq. 2. Blue dashed lines are from a global fit to resolution convolved single modes with dispersion as listed in Eq. 6.

SPINS was used to probe the inter-chain interactions with a final energy $E_f = 2.5$ meV resulting in a measured full width at half maximum (FWHM) elastic energy resolution of $\delta\hbar\omega = 50 \mu\text{eV}$. Magnetic field dependent measurements at SPINS were measured with $E_f = 2.8$ meV. This resulted in an average FWHM energy resolution of $\delta E = 0.12$ meV and FWHM \mathbf{Q} resolution along the projected chain direction of $\delta Q_{\parallel} = 0.03 \text{ \AA}^{-1}$.⁷³ Magnetic field dependent measurements at the DCS were performed with four instrument configurations as listed in Table I. Details concerning the spectrometer configurations for the SPINS and DCS instrument are discussed in the Supplemental Information.⁷¹

IV. EXPERIMENTAL RESULTS

A. Interchain coupling

To determine the spin Hamiltonian for CN a zero field inelastic neutron scattering experiment was conducted on the SPINS triple-axis spectrometer at NIST with the sample oriented in the (hkh) zone. Figure 2 shows three constant \mathbf{Q} scans for $T = 1.25$ K, which is well above the critical temperature for magnetic ordering. Two modes are observed at $(1.25, 0.5, 1.25)$ (Fig. 2(a)), while only a single mode is observed at $(1.5, 0, 1.5)$ and $(1, 0, 1)$, (Figs. 2(b) and (c)). The peaks in Figs. 2(b) and (c) are at similar energy transfers as the peaks in Fig. 2(a).

Configuration	E_i (meV)	f (Hz)	resolution mode	$\hbar\omega_{\min}$ (meV)	$\hbar\omega_{\max}$ (meV)	$\text{FWHM}\delta\hbar\omega$ (μeV)	$ \mathbf{Q}_{\min} $ (\AA^{-1})	$ \mathbf{Q}_{\max} $ (\AA^{-1})
1	2.10	333.3	medium	-0.12	1.10	28	0.18	1.79
2	1.90	333.3	low	-0.15	0.95	50	0.17	1.70
3	1.132	266.8	low	-0.04	0.57	29	0.13	1.31
4	1.90	279.95	low	-0.15	1.06	57	0.17	1.70

TABLE I: Experimental configurations for measurements using the DCS at NIST. The frequency of neutron pulses at the sample is f . The resolution mode is determined by the widths of the chopper slots phased to transmit the beam; see Ref. 72. Minimum and maximum energy transfers correspond to first and last channels in the time-of-flight histogram. The FWHM $\delta\hbar\omega$ is the energy resolution at the elastic position. The minimum and maximum wave vector transfers are also given. The configuration number is referenced in figure captions.

Furthermore, all of the peaks in Fig. 2 are located at the top or the bottom of the band of magnetic excitations previously measured for CN.⁶¹ Based on a Gaussian peak approximation, the mode energies for the data in Fig. 2 are $\hbar\omega = 0.39(2)$ meV and $0.49(2)$ meV, and the energies of the peaks in Figs. 2(b) and (c) are $\hbar\omega = 0.49(2)$ meV and $0.40(2)$ meV respectively.⁷⁴

For the $[1 \pm 1 1]$ alternating chain model, there are two distinct modes propagating in the (hkh) plane. Here denoted A and B, these correspond to the two directions of alternating chains, $[111]$ and $[\bar{1}\bar{1}\bar{1}]$. Considering interactions, J'_2 , between $[1 \pm 1 1]$ chains gives the following dispersion relations.

$$\begin{aligned}
\epsilon_A(\mathbf{Q}) &= J_1 - \frac{J_2}{2} \cos(\pi(h+k+l)) \\
&\quad - \frac{J'_2}{2} \cos(\pi(h-k+l)) \\
\epsilon_B(\mathbf{Q}) &= J_1 - \frac{J_2}{2} \cos(\pi(h-k+l)) \\
&\quad - \frac{J'_2}{2} \cos(\pi(h+k+l)).
\end{aligned} \tag{6}$$

Note that if the primary chain direction is along $[111]$ ($[\bar{1}\bar{1}\bar{1}]$), then interchain interactions are along the $[\bar{1}\bar{1}\bar{1}]$ ($[111]$) direction as shown in Figs. 1(d) and (e). The difference in mode energies for the wave-vectors in Figs. 2(b) and (c) is $J_2 + J'_2$, and the difference in energies of the excitations in Fig. 2(a) is $J_2 - J'_2$. The measured difference in the energies of the modes in Figs. 2(b) and (c) and the modes in Fig. 2(a) are $0.09(2)$ and $0.09(2)$ meV respectively. From Eq. 6, the difference in these values limits the magnitude of inter-dimer interactions perpendicular to the primary chain direction to $2\tilde{J}'_2 = -0.01(2)$ meV. As described earlier in regards to the \tilde{J}_2 , the “ \sim ” indicates that we are reporting $T = 1.25$ K measurements so the inferred energies can be expected to be renormalized with respect to actual exchange constants.⁶¹ The mean difference in peak locations in Fig. 2 corresponds to inter-dimer exchange $\tilde{J}_2 = 0.09(2)$ meV at $T = 1.25$ K. Our determined values of \tilde{J}'_2 are consistent with zero, and therefore it is appropriate to describe CN as well isolated alternating chains that run in the $[1 1 1]$ and $[1 \bar{1} \bar{1}]$ directions on adjacent crystal planes spanned by \mathbf{b} and $\mathbf{a} + \mathbf{b}$.

We also performed global fits to these data using the dispersion relations in Eq. 6 and a dimer structure factor convolved with the instrumental resolution function (dashed lines in Fig. 2). The results are consistent with the simplified fitting analysis: $\tilde{J}'_2 = -0.01(2)$ meV, $J_1 = 0.44(2)$ meV and $\tilde{J}_2 = 0.08(2)$ meV for $T = 1.25$ K. A similar global fit with $\tilde{J}'_2 \equiv 0$ and a dimer structure factor yields $J_1 = 0.44(2)$ meV and $\tilde{J}_2 = 0.09(2)$ meV, solid lines in Fig. 2. The values for J_1 and \tilde{J}_2 determined for $T = 1.25$ K are consistent with previous inelastic neutron scattering measurements taking into account the temperature dependent bandwidth normalization factor.⁶¹

In the following, our analysis of the field dependent excitation spectrum in CN only includes the J_1 and J_2 exchange parameters. Although we make structural arguments above concerning the correct inter-dimer linkage, the current experiment does not identify which of the two types of linkage creates the alternating chains, only that one of them corresponds to the alternating chain and the other is very weak.

B. Field Dependent Magnetic Excitations

1. Triple-axis measurements

Figures 3(a)-(c) show $\tilde{q} - \hbar\omega$ maps of neutron scattering intensity at $T = 1.8$ K measured for $\mu_0 H = 0$, $\mu_0 H = 1.5$ T, and $\mu_0 H = 7$ T on the SPINS triple-axis spectrometer. In zero magnetic field, there is a single resonant mode with an energy gap of $\Delta \approx 0.35$ meV and a bandwidth of approximately $90 \mu\text{eV}$. The spectrum appears broadened in comparison to prior zero-field measurements.⁶¹ This is due to both the elevated temperature and a factor of five coarser energy resolution compared to the prior backscattering measurements. The periodicity in \tilde{q} of this mode is consistent with Eq. 6.

At $\mu_0 H = 1.5$ T, Fig. 3(b) shows three resolution limited modes. Two of these are clearly seen at $\hbar\omega \approx 0.4$ and 0.6 meV. The third is located in the vicinity of 0.2 meV though it is only apparent through increased scattering intensity in this energy range at field over the corresponding zero-field data. The dispersion of these modes is indistinguishable from the zero field dispersion, $\epsilon_0(\mathbf{Q})$,

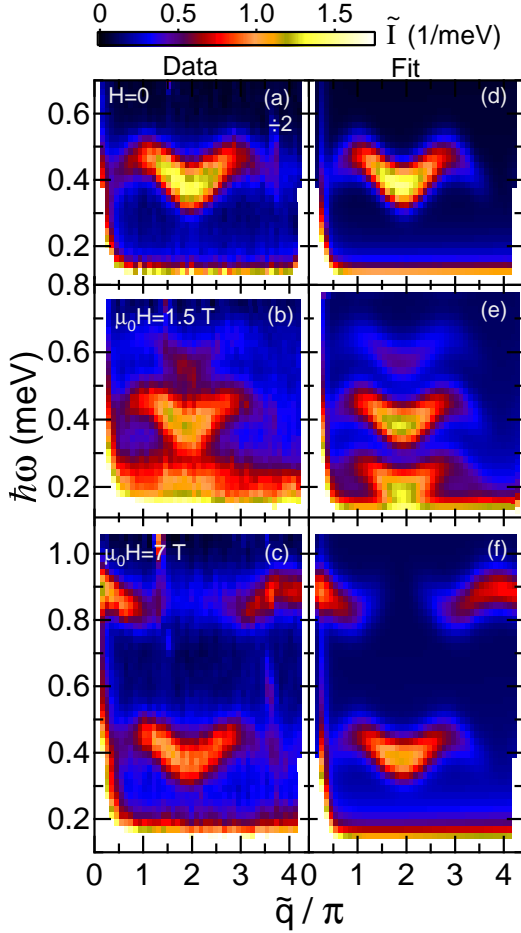


FIG. 3: (color online) Normalized scattering intensity, $\tilde{I}(\tilde{q}, \hbar\omega)$, for CN at $T = 1.8$ K for (a) $H = 0$, (b) $\mu_0 H = 1.5$ T, and (c) $\mu_0 H = 7$ T from triple-axis measurements. Zero field results are shown as half the measured/calculated intensity. Panels (d)-(f) are the results of fits described in the text.

with splitting given by the Zeeman term:

$$\epsilon(\mathbf{Q}) = \epsilon_0(\mathbf{Q}) - g\mu_B\mu_0 H S_z. \quad (7)$$

For CN in the $(h0l)$ scattering plane with $J'_2 \approx 0$

$$\epsilon_0(\mathbf{Q}) = J_1 - \frac{J_2}{2} \cos(\pi(h+l)), \quad (8)$$

from Eq. 6. The splitting of the spectrum confirms the singlet-triplet nature of the magnetic excitations in CN as described below in Fig. 11 for $H < H_{c1}$.

Above the upper critical field, $\mu_0 H_{c2} = 4.2$ T, CN is fully magnetized at sufficiently low T . Figure 3(c) shows the $\tilde{q} - \hbar\omega$ dependent scattering intensity for $\mu_0 H = 7$ T where two modes are observed. The lower energy mode has similar intensity modulation and dispersion versus \tilde{q} as in zero field. In contrast, the \tilde{q} dependence of the upper mode is out of phase with the lower mode.

Data for the intermediate field regime, $\mu_0 H = 3.55$ T, is shown in Fig. 4(a). The mode at $\hbar\omega \approx 0.4$ meV is

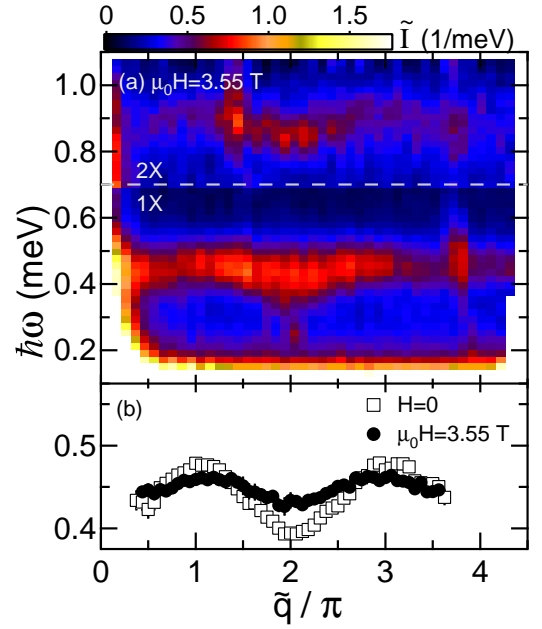


FIG. 4: (color online) (a) Normalized scattering intensity, $\tilde{I}(\tilde{q}, \hbar\omega)$, for CN at $T = 1.8$ K and $\mu_0 H = 3.55$ T from triple-axis measurements. For clarity, data above the white dashed line ($\hbar\omega > 0.7$ meV) have been multiplied by a factor of two. (b) Peak positions for $\hbar\omega \approx 0.45$ meV for constant \tilde{q} scans through $H = 0$ and $\mu_0 H = 3.55$ T spectra at $T = 1.8$ K.

broadened compared to data collected below the lower critical field. The upper mode of the triplet, although reduced in intensity, has shifted to higher energies with unchanged dispersion and intensity modulation. Fig. 4(b) shows fitted peak positions for the zero field and $\mu_0 H = 3.55$ T mode, illustrating the change in dispersion at intermediate fields.

For a parametric overview of the field dependence, constant \tilde{q} scans were measured at $\tilde{q} = n\pi$ at integer n for a range of fields covering all portions of the magnetic phase diagram at $T = 1.8$ K. For select values of the applied magnetic field, these data are shown in Fig. 5 for $\tilde{q} = 2\pi$ and Fig. 6 for $\tilde{q} = \pi$.

In zero field, there is a single resolution limited mode at the bottom and top of the magnetic band. For magnetic fields up to 2 T, all three of the Zeeman split triplet modes are visible. However, for larger magnetic fields, the lowest energy excitation of the triplet is obscured by incoherent elastic nuclear scattering. Above the lower critical field, $\mu_0 H_{c1} = 2.8$ T, a mode persists at $\hbar\omega \approx 0.45$ meV, in addition to the higher energy mode which continues to rise in energy. An overall broadening of the $\hbar\omega \approx 0.45$ meV mode is apparent in the intermediate field regime. Comparing the lineshapes for $\tilde{q} = 2\pi$ and $\tilde{q} = \pi$, the $\tilde{q} = 2\pi$ scans are consistently broader than the $\tilde{q} = \pi$ scans. This is also seen in the \tilde{q} dependent spectrum measured at $\mu_0 H = 3.55$ T in Fig. 4(a). At $\mu_0 H = 4.2$ T, the peak at $\hbar\omega \approx 0.45$ meV becomes further broadened, and at $\mu_0 H = 5$ T a second peak is

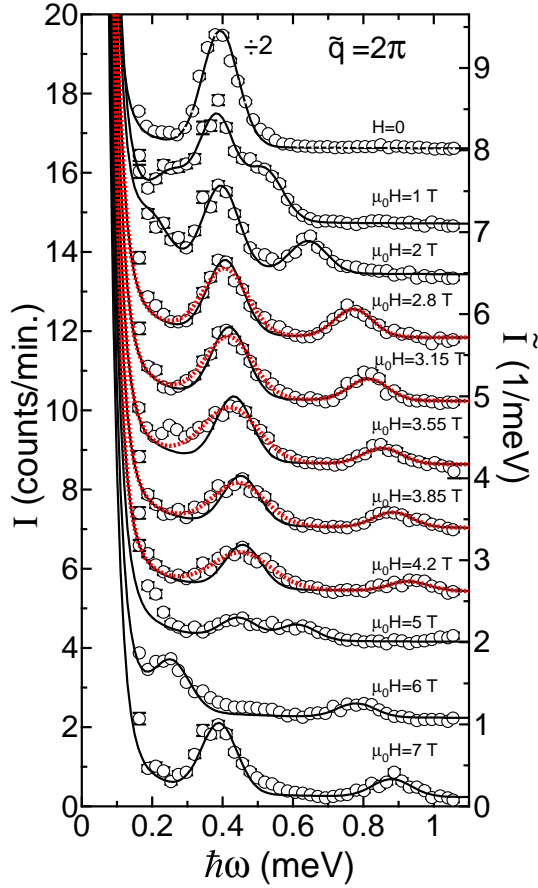


FIG. 5: (color online) Normalized scattering intensity, $\tilde{I}(\tilde{q} = 2\pi, \hbar\omega)$, for CN at $T = 1.8$ K for magnetic fields $0 \leq \mu_0 H \leq 7$ T from triple-axis measurements. Results have been offset vertically for presentation. Solid lines are fits to resolution limited Gaussians as described in the text. Dotted red lines for $2.8 \leq \mu_0 H \leq 4.2$ T are fits to resolution limited Gaussians with an additional FWHM added in quadrature as described in the text. Data below $\hbar\omega = 0.15$ meV, which are dominated by incoherent elastic nuclear scattering, have been omitted for clarity.

apparent at $\hbar\omega \approx 0.6$ meV. At $\mu_0 H = 6$ and $\mu_0 H = 7$ T, the two modes in Figs. 5 and 6 correspond to those depicted in Fig. 3(c).

2. Time-of-flight measurements

Excitations in the intermediate field range, $H_{c1} \leq H \leq H_{c2}$, were examined with the higher resolution time-of-flight instrument. Figure 7 shows data for a range of magnetic fields collected at $T = 0.224$ K, which exceeds the maximum Néel temperature. The particular magnetic fields chosen correspond to the rational fractions of the low-temperature saturation magnetization indicated within the figure.⁷⁵

We first compare the zero-field data (Fig. 7(a)) to the SPINS data. The modes are sharper due to the en-

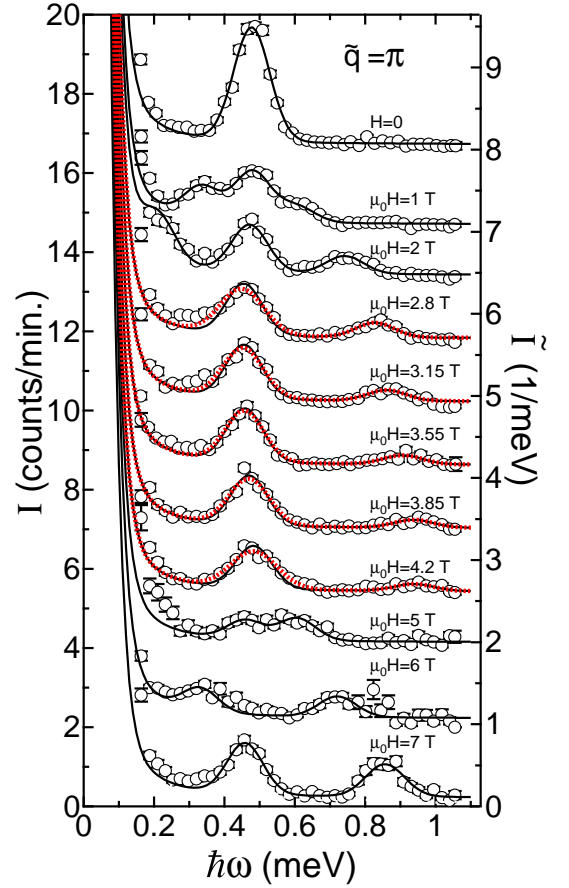


FIG. 6: (color online) Normalized scattering intensity, $\tilde{I}(\tilde{q} = \pi, \hbar\omega)$, for CN at $T = 1.8$ K for magnetic fields $0 \leq \mu_0 H \leq 7$ T from triple-axis measurements. Results have been offset vertically for presentation. Solid lines are fits to resolution limited Gaussians as described in the text. Dotted red lines for $2.8 \leq \mu_0 H \leq 4.2$ T are fits to resolution limited Gaussians with an additional FWHM added in quadrature as described in the text. Data below $\hbar\omega = 0.15$ meV, which are dominated by incoherent elastic nuclear scattering, have been omitted for clarity.

hanced energy resolution. This, as well as the lower temperature⁶¹ leads to the increased peak scattering intensity. For the Shastry-Sutherland spin dimer system $\text{SrCu}_2(\text{BO}_3)_2$ the singlet triplet transition near 3 meV is split by 0.4 meV or 13%. For comparison Fig. 7(a) places an upper limit of 0.02 meV or 5% on any zero-field peak splitting. This indicates the Heisenberg approximation to intra-dimer spin interactions is excellent for CN.

At $\mu_0 H = \mu_0 H_{c1} = 2.82$ T, Fig. 7(b), the three modes of the zero field triplet are split. The central, $S_z = 0$, mode is at the same location as in zero-field. The $S_z = \pm 1$ modes are less intense and Zeeman split to higher and lower values of $\hbar\omega$. The $S_z = -1$ mode lies at $\hbar\omega \approx 0.75$ meV, while the $S_z = 1$ mode is manifest in increased scattering intensity near the incoherent elastic line.

At higher fields, the upper mode continues to increase in energy while the central mode at $\hbar\omega \approx 0.45$ meV

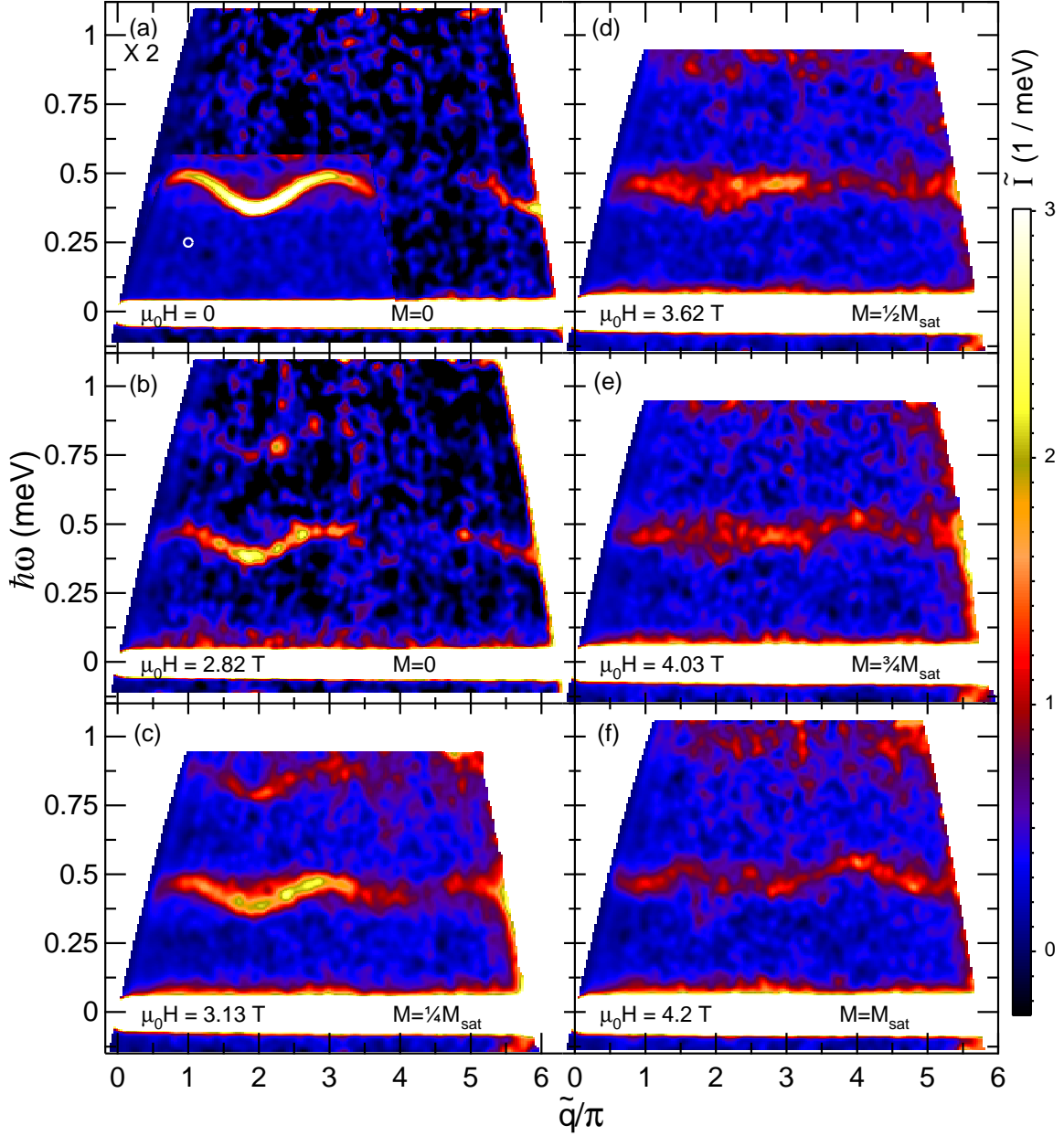


FIG. 7: (color online) Normalized scattering intensity, $\tilde{I}(\tilde{q}, \hbar\omega)$ for CN at $T = 0.224$ K for (a) $H = 0$, (b) $\mu_0 H = 2.82$ T, (c) $\mu_0 H = 3.13$ T, (d) $\mu_0 H = 3.62$ T, (e) $\mu_0 H = 4.03$ T, and (f) $\mu_0 H = 4.2$ T from time-of-flight measurements. The data in (a) are shown at half the intensity scale. The trapezoidal area in (a) was covered by the high resolution Config. 3 while the surrounding area was measured with Config. 1. Data in (b) were measured with Config. 1. Data in (c), (d), and (e) were measured with Config. 2. Data in (f) were measured with Config. 4. False color images of the scattering intensity were produced by binning the data onto an orthogonal grid in $\hbar\omega$ and \tilde{q} . The data were then convolved with a normalized Gaussian with FWHM principal axes of $\hbar\omega = 31 \mu\text{eV}$ and $\tilde{q}/\pi = 0.125$. The corresponding FWHM ellipsoid is shown in (a), and approximates the average resolution ellipsoid.

changes its \tilde{q} -dependence and the intensity distribution moves to slightly higher values of $\hbar\omega$. At $\mu_0 H = \mu_0 H_{c2} = 4.2$ T, Fig. 7(f), the \tilde{q} dependence of both the peak position and intensity associated with the central mode – considered as a function of \tilde{q} – have changed phase.

For an overview of the magnetic density of states versus

field, the wave-vector integrated intensity ($\frac{1}{2}\pi < \tilde{q} < \frac{7}{2}\pi$) is shown in Fig. 8. In zero field, there are peaks associated with the Van-Hove singularities at the top and bottom of the magnetic excitation spectrum. At the lower critical field, $\mu_0 H_{c1} = 2.82$ T, these singularities become less pronounced and the upper $S_z = -1$ band appears as a

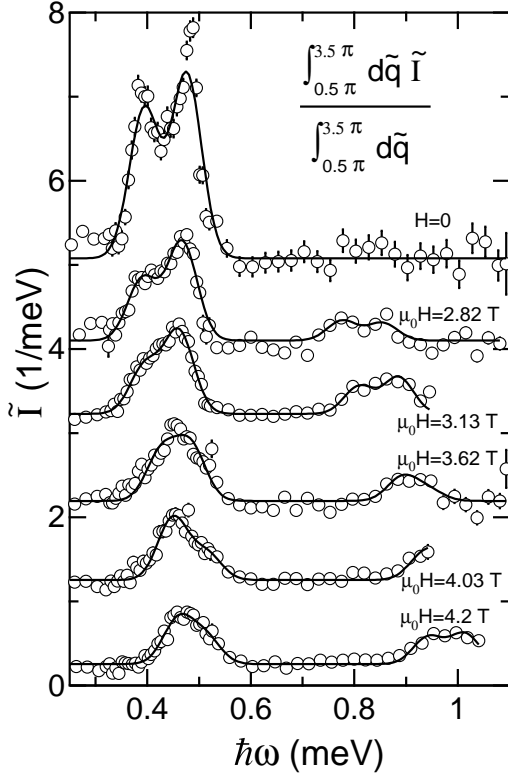


FIG. 8: Wave-vector averaged magnetic scattering intensity for CN at $T = 0.224$ K as a function of $\hbar\omega$ and applied magnetic field from time-of-flight measurements. The average covered the range from $\tilde{q} = \frac{1}{2}\pi$ to $\tilde{q} = \frac{7}{2}\pi$. Data for different fields are offset vertically for presentation purposes. Solid lines are the result of gaussian fits as described in the text. The $H = 0$ and $\mu_0 H = 2.82$ T data were combined from measurements made in Configs. 1 and 3. The $\mu_0 H = 3.13$ T and $\mu_0 H = 4.03$ T data were acquired using Config 2. The $\mu_0 H = 3.62$ T data were combined from measurements made in Configs. 1, 2, and 3. The $\mu_0 H = 4.2$ T data are from measurements made in Config. 4.

rounded double peak at $\hbar\omega \approx 0.8$ meV. With increasing field both bands shift to higher energies, with a greater rate for the higher energy $S_z = -1$ band.

Figures 9 and 10 show constant $\tilde{q} = 2\pi$ and $\tilde{q} = \pi$ scans extracted from the data shown in Fig. 7. In zero field at $\tilde{q} = 2\pi$, there is a single peak at $\hbar\omega \approx 0.4$ meV corresponding to crossing the bottom of the magnetic band. As the magnetic field is increased, this peak broadens, loses intensity, and moves to higher values of energy transfer. An analogous development with field is observed for the upper branch of the Zeeman split triplet, which broadens and shifts from $\hbar\omega = 0.75$ meV to $\hbar\omega = 0.95$ meV upon raising the field from the lower critical field $\mu_0 H = \mu_0 H_{c1} = 2.8$ T to the upper critical field $\mu_0 H = \mu_0 H_{c2} = 4.2$ T.

For $\tilde{q} = \pi$ (Fig. 10) the single peak at $\hbar\omega \approx 0.45$ meV corresponds to crossing the top of the magnetic band in zero field. With increasing field, this peak broadens, and also shifts slightly to lower energy transfer. Just as for

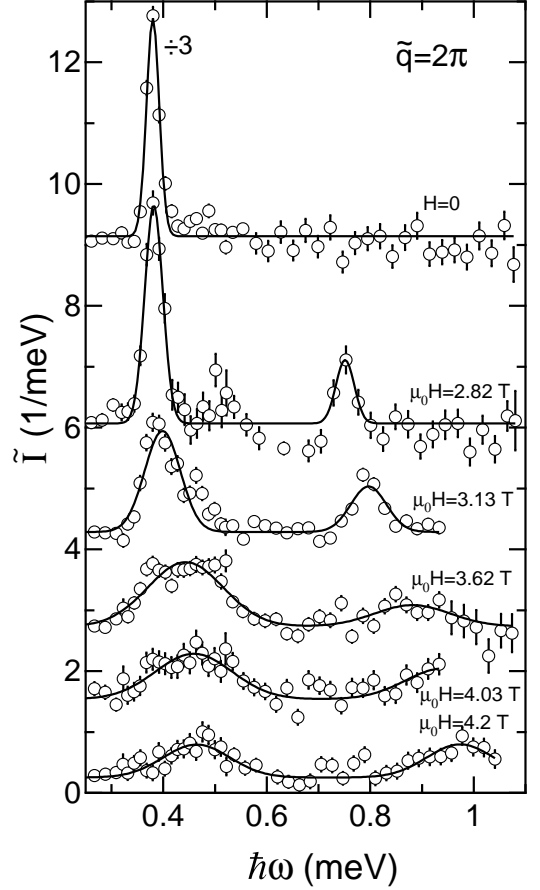


FIG. 9: Constant $\tilde{q} = 2\pi$ scattering intensity ($\delta\tilde{q} = 0.13\pi$) as a function of $\hbar\omega$ and applied magnetic field for CN at $T = 0.224$ K from time-of-flight measurements. The zero-field data has been reduced in intensity by a factor of three for presentation purposes. Solid lines are fits to Gaussians as described in the text. Results have been offset vertically for presentation. The $H = 0$ and $\mu_0 H = 2.82$ T data were combined from measurements made in Configs. 1 and 3. The $\mu_0 H = 3.13$ T and $\mu_0 H = 4.03$ T data are from Config 2. The $\mu_0 H = 3.62$ T data were combined from measurements made in Configs. 1, 2, and 3. The $\mu_0 H = 4.2$ T data are from measurements made in Config. 4.

$\tilde{q} = 2\pi$, the higher energy $S_z = -1$ branch becomes visible for $H > H_{c1}$ and shifts to higher $\hbar\omega$ with increasing fields.

V. ANALYSIS AND DISCUSSION

While the resolution employed here is adequate to resolve the dispersion of gapped excitations, quasi-elastic features in the intermediate field regime are not well separated from the elastic nuclear scattering.⁷⁶ Our discussion therefore focuses on gapped excitations, first covering fields below H_{c1} and above H_{c2} , then the intermediate field regime. For context we start by presenting the re-

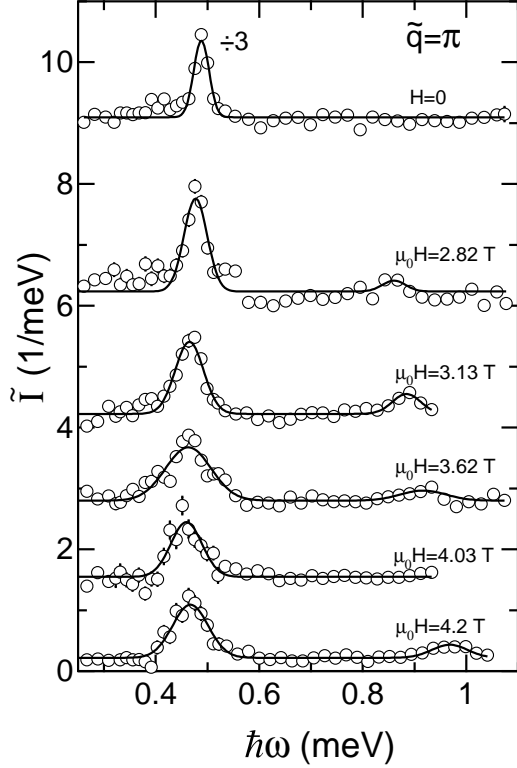


FIG. 10: Constant $\tilde{q} = \pi$ scattering intensity ($\delta\tilde{q} = 0.13\pi$) as a function of $\hbar\omega$ and applied magnetic field for CN at $T = 0.224$ K from time-of-flight measurements. The zero-field data has been reduced in intensity by a factor of three. Solid lines are fits to Gaussians as described in the text. Results have been offset vertically for clarity. The $H = 0$ and $\mu_0 H = 2.82$ T data were combined from measurements made in Configs. 1 and 3. The $\mu_0 H = 3.13$ T and $\mu_0 H = 4.03$ T data are from Config 2. The $\mu_0 H = 3.62$ T data were combined from measurements made in Configs. 1, 2, and 3. The $\mu_0 H = 4.2$ T data are from measurements made in Config. 4.

sults of exact diagonalization calculations for finite length alternating spin chains in an applied magnetic field.

A. Exact Diagonalization

Exact diagonalization calculations for finite length alternating spin-1/2 chains are instructive to understand the magnetic field dependent phase diagram and excitation spectra of CN. We computed the eigenvalues of the Hamiltonian of Eq. 1 with $\alpha = \frac{1}{4}$ for even-length chains up to $N = 16$ spins, using code we previously developed for the analysis of a gapped 1D quantum spin system.⁷⁷ The computations were carried out at $H = 0$ using standard matrix diagonalization routines,⁷⁸ and we used the wave vector \tilde{q} (via periodic boundary conditions) as well as the z -component of the total spin S_z to reduce the sizes of the non-zero subspaces of the Hamiltonian matrix to be diagonalized. The total spin quantum number

S for each state was inferred by matching the energies of states with $S_z < N/2$ with those at higher S_z .

Figure 11 gives results of these computations for $N = 6$ and $N = 16$, and shows the evolution with magnetic field of the lowest energy multiplet at each S , the $\tilde{q} = 0$ excitation spectra at $T = 0$, and the ground state magnetization. In Figs. 11(a) and (b), we label the allowed $T = 0$, dipole-active transitions from (i) to (vi) based upon their respective values of S and S_z .⁸¹

The excitations in zero field are between a non-magnetic singlet ground state and a magnetic triplet ($S = 1$). On application of an external magnetic field, the degeneracy of the three excitations shown in Fig. 11(a), (b), (d) and (e) is lifted through Zeeman splitting of the triplet. For larger magnetic fields, there exists a quantum critical point (QCP), H_{c1} , where the spin-gap is closed and magnetization develops as a series of higher- S_z states successively becomes the ground state. In the intermediate field regime above H_{c1} , a total of five non-trivial transitions are allowed (see Fig. 11(a) and (b)).

As shown most clearly for our longest chain calculations ($N = 16$) in the right panels of Fig. 11, the field-dependence of the finite energy excitations in this regime becomes linear with an intercept at the origin. There is also a set of low-energy excitations, which in the $N \rightarrow \infty$ limit should yield a gapless regime, and inter-chain interactions are expected to produce a Néel-like ground state. At higher fields there is a second QCP, H_{c2} , beyond which the system is fully polarized in a forced ferromagnetic state with two transverse spin-wave modes. The calculated magnetization in these three field ranges is consistent with the measured low-temperature magnetization for CN (Fig. 11(f)). In the following we shall discuss the experimental observations in light of these exact diagonalization results.

B. $H < H_{c1}$ and $H > H_{c2}$

The intensity modulation of the modes in Fig 3(a)-(c) is not commensurate with the crystal lattice. This can be understood by examining the dynamic spin correlation function for a dimer antiferromagnet in conjunction with the SMA in the absence of anisotropy.^{79,80} The SMA has been used with success to describe the zero field modes in copper nitrate such that the dynamic spin correlation function becomes⁶¹

$$S(\mathbf{Q}, \hbar\omega) = \frac{J_1 \langle \mathbf{S}_0 \cdot \mathbf{S}_{\mathbf{d}_1} \rangle}{3\epsilon(\mathbf{Q})} [1 - \cos(\mathbf{Q} \cdot \mathbf{d}_1)] \times \delta(\hbar\omega - \epsilon(\mathbf{Q})), \quad (9)$$

where \mathbf{d}_1 is the intra-dimer vector. This expression describes the scattering associated with singlet to triplet transitions. In the magnetized state however, transitions can occur between states that originate from the zero field triplet. For an isolated spin pair these carry a structure factor proportional to $(1 + \cos(\mathbf{Q} \cdot \mathbf{d}_1))$.

For the case of a general spin cluster, the $[1 - \cos(\mathbf{Q} \cdot \mathbf{d}_1)]$ structure factor in Eq. 9 needs to be modified to ac-

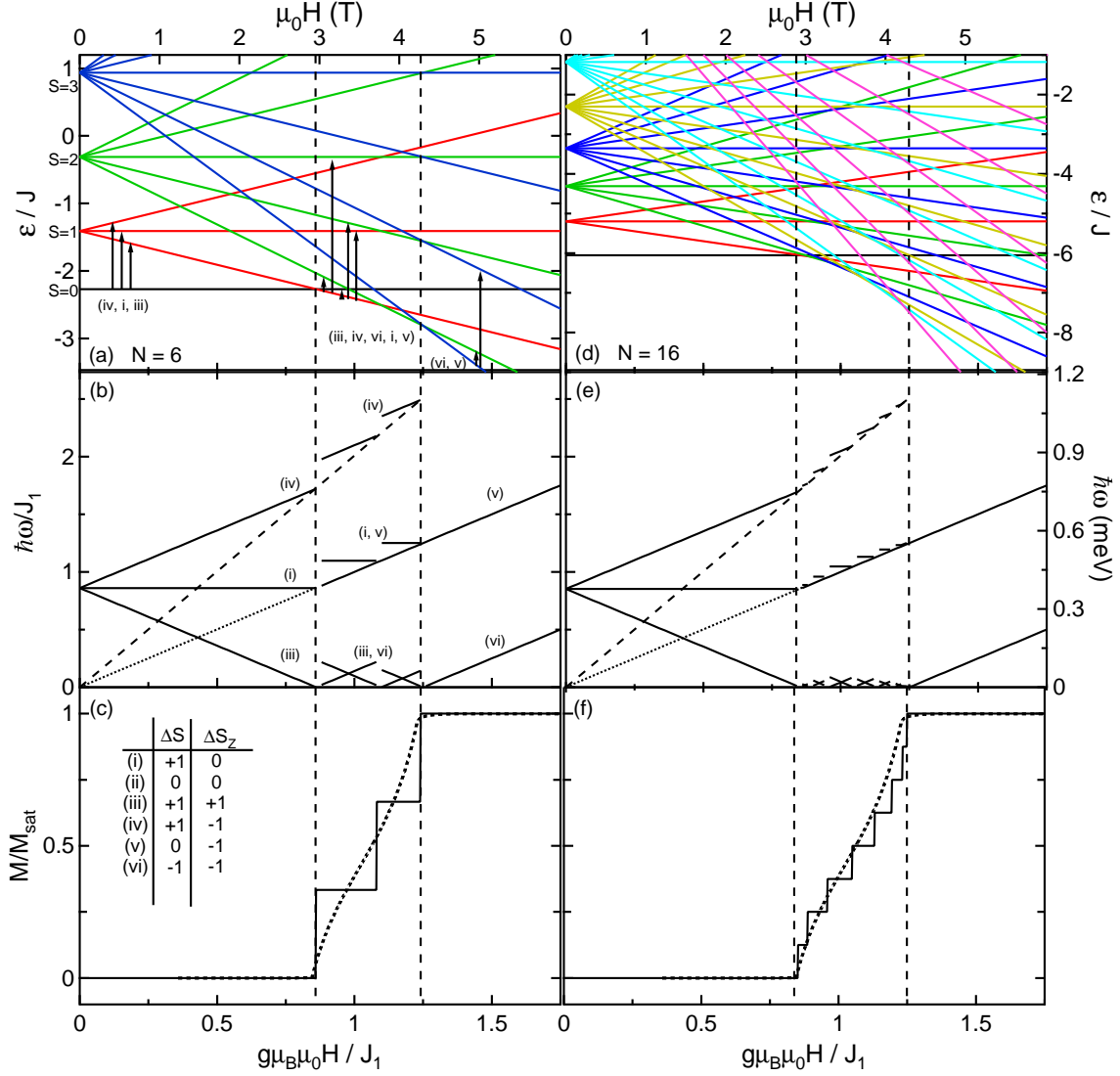


FIG. 11: (color online) Zero temperature exact diagonalization calculations for $N = 6$ (a-c) and $N = 16$ (d-f) $\alpha = 0.25$ alternating antiferromagnetic Heisenberg spin chains as a function of applied magnetic field at zero wave-vector. Panels (a) and (d) are the calculated field-dependent energy levels. Only the lowest-energy multiplet at each total spin S is shown. States illustrated in the same color in both (a) and (b) have the same S . Panel (d) also shows states with $S = 4$ (olive), $S = 5$ (light blue), and $S = 8$ (pink). Multiplets with $S = 6$ and $S = 7$ are omitted for clarity. Panels (b) and (e) are the lowest energy dipole active transitions. Dotted and dashed lines are extrapolations of the field dependence of excitations in the intermediate field regime. Panels (c) and (f) are the calculated magnetization based on the total moment of the ground state. The dotted line in these panels is the measured $T = 0.1$ K field dependent magnetization of CN.⁷⁵ Panels (a) and (b) also identify transitions in the basis of total spin S and the z -component of spin S_z using lower case Roman numerals (i) through (vi) as listed in the table in panel (c) and described in Ref. 81. Reduced units are used for the bottom axes, and units of applied magnetic field for the case of $J_1 \approx 0.44$ are used for the top axes.

count for the initial S and final S' spin quantum numbers of the excitation.^{82,83} We therefore consider the following generalization of Eq. 9 in analyzing the magnetic excitations in a field,

$$S(\mathbf{Q}, \hbar\omega) = \frac{J_1 \langle \mathbf{S}_0 \cdot \mathbf{S}_{\mathbf{d}_1} \rangle}{3\epsilon(\mathbf{Q})} [1 + (-1)^{S-S'} \cos(\mathbf{Q} \cdot \mathbf{d}_1)] \times \delta(\hbar\omega - \epsilon(\mathbf{Q})). \quad (10)$$

Here S and S' refer to the initial and final spin state of the transition labeled in accordance with the non-interacting limit. Equation 10 retains the oscillator strength for each mode while introducing the anticipated dispersion from perturbation theory. The expression is appropriate in the limit of $\alpha = J_2/J_1 \rightarrow 0$ and for the forced ferromagnetic state as described shortly.

For $H < H_{c1}$, the low energy states of the antiferro-

magnetic alternating chain are the $S = 0$ singlet ground state and the $S = 1$ triplet excited state (Fig. 11). The cosine term in Eq. 10 has a negative prefactor for these transitions. Figure 3(d) is a fit to the zero field data based on equations 7 through 10 including an incoherent elastic background. This model spectrum has three fitting parameters: two exchange constants, J_1 and J_2 , and an overall multiplicative prefactor. The model accounts well for the scattering intensity and dispersion with $J_1 = 0.44(4)$ meV and $\tilde{J}_2 = 0.09(4)$ meV at $T = 1.8$ K. The values of intra- and interdimer exchange are consistent with the values obtained from measurements at similar temperatures described earlier and with the temperature dependent renormalization of the bandwidth determined by Xu *et al.*⁶¹

To describe the data in Fig. 3(b) we employed the Zeeman split dispersion as described in Eq. 7 with $\epsilon_0(\mathbf{Q})$ given by Eq. 8 along with Eqs. 10 and 4. Considering the potential for an anisotropic g -factor, prefactors for the $S_z \neq 0$ components were allowed to vary and this resulted in the excellent fit shown in Fig. 3(e). The corresponding values of $J_1 = 0.43(4)$ meV, $\tilde{J}_2 = 0.07(4)$ meV are consistent with the zero field fits. The intensities of the lower and upper modes respectively were found to be 39(1)% and 47(2)% of the $S_z = 0$ corresponding on average to $\delta g = 0.17(1)$, which is consistent with $g = \sqrt{g_b^2 + g_\perp^2} = 2.22$ derived from bulk measurements.⁸⁴

For $H > H_{c2}$, the \tilde{q} dependence of the ω -integrated intensity for the lower and upper modes can be understood in terms of Eqs. 10 and 4 in conjunction with the transitions illustrated in Fig. 11. From our finite chain calculations, the change in total spin from the ground state to the lowest excited state for the ferromagnetic phase is odd, $\Delta S = -1$, while the change in total spin to the next highest excited state is even, $\Delta S = 0$. This leads to the phase difference in the intensities of the two modes in the $\mu_0 H = 7$ T spectrum through the $(-1)^{S-S'}$ term in Eq. 10.

For a simulation of the high field data we use the Holstein-Primakoff spinwave calculation, to calculate the dispersion of a 1d alternating ferromagnet as

$$\epsilon_{FSW}(\mathbf{Q}) = g\mu_B(H - H_{c2}) + \frac{1}{2}(J_1 + J_2) \pm \frac{1}{2}\sqrt{J_1^2 + J_2^2 + 2J_1J_2\cos(\mathbf{Q} \cdot \mathbf{u})}. \quad (11)$$

where S is the spin of the magnetic ions, J_1 and J_2 are the intra- and inter-dimer exchange and \mathbf{u} is the inter-dimer vector. Here the applied magnetic field term has been offset by the upper critical field. The dispersion in Eq. 11 is identical to that found via a bosonization representation of the excitations of the alternating ferromagnetic chain.⁸⁵

The data in Fig. 3(c) were fit using Eqs. 10 - 11 and a comparison between model and data is shown in Fig. 3(f). The best fit exchange parameters $J_1 = 0.46(4)$ meV, $\tilde{J}_2 = 0.07(4)$ meV are consistent with measurements from the singlet ground state. The fit included a multiplicative

scale factor corresponding to the ratio of intensities of the upper band to the lower band to account for the g -factor.

CN is thus well described by the alternating chain model for magnetic fields below the lower critical field and above the upper critical field. The fitted exchange constants found for $H < H_{c1}$ and $H > H_{c2}$ are similar even though the non-negligible temperature of the measurements may affect the results near the critical fields. The \tilde{q} dependence of the low energy modes found in the exact diagonalization calculations is consistent the measured spectra below H_{c1} and above H_{c2} .

For an overview of the field dependence of the excitation spectrum we fit the constant $\tilde{q} = \pi$ and $\tilde{q} = 2\pi$ of Figs. 5 and 6 to sums of resolution limited Gaussians. The $\hbar\omega$ -dependent background determined from analysis of the full data set of Fig. 3 was used. The field-dependent peak positions for $\tilde{q} = 2\pi$ are shown in Fig. 12. We plot the field dependence of the calculated $H < H_{c1}$ and $H > H_{c2}$ excitations at $\tilde{q} = 2\pi$ as solid lines in Fig. 12 with $g = \sqrt{g_b^2 + g_\perp^2} = 2.22$ ⁸⁴ using the low-temperature values of the exchange constants.⁶¹ There is very good agreement between the measured and calculated field dependence.

While the modes observed for $\mu_0 H \geq 6$ T data are consistent with Eq. 11, spectra for fields between $\mu_0 H_{c2}$ and $\mu_0 H = 5$ T are not. There are two resolution limited modes (represented by inverted and upright triangles in Fig. 12), but they are not separated by the same energy as for the $\mu_0 H \geq 6$ T data. Immediately above $\mu_0 H_{c2}$, the higher energy mode is consistent with theory, but the lower energy modes occur at elevated values of $\hbar\omega$. This can be understood by considering the temperatures used in the measurements. Just above $\mu_0 H_{c2}$, low energy states are thermally populated due to the relatively high temperature of the measurement: $T = 1.4$ K. The additional excitations observed for $\mu_0 H = 4.6$ and 5 T are due to scattering from the excited states rather than the ground state, and we label them as $\Delta S=1$, and $\Delta S_z=0$. The energy of the zero temperature lowest energy excitation out of the ground state then corresponds to the difference in energies of the two modes observed. These values are plotted as crosses inside of open squares in Fig. 12, and are consistent with the field dependent ferromagnetic dispersion above H_{c2} .

C. Intermediate fields, $H_{c1} < H < H_{c2}$

While the magnetization is constant for fields below H_{c1} and above H_{c2} it varies continuously with field between these critical values. The implication is a gapless excitation spectrum that cannot support truly long-lived magnetic excitations. Near the elastic line a spectrum resembling that of the gapless spin-1/2 chain is anticipated but the instrumentation utilized here cannot resolve these features from the intense incoherent elastic nuclear scattering.

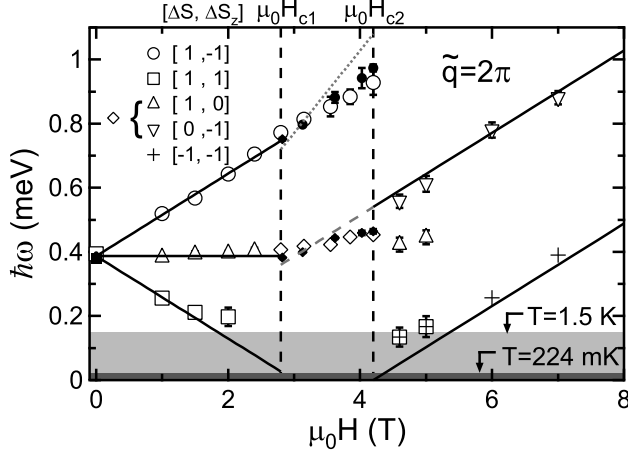


FIG. 12: Fitted peak position in the $\tilde{q} = \pi$ magnetic neutron scattering spectrum of CN as a function of applied magnetic field for $T = 1.8$ K (open symbols, triple-axis measurements) and $T = 0.224$ K (closed symbols, time-of-flight measurements). The raw data and fits are shown in Figs. 5 and 9. $\mu_0 H_{c1} = 2.8$ T and $\mu_0 H_{c2} = 4.2$ T are indicated by dashed vertical lines. For excitations near $\hbar\omega \approx 0.45$ meV at intermediate fields, fits were performed with an additional FWHM parameter 2Γ added in quadrature to the resolution width. The time-of-flight constant \tilde{q} scans in the intermediate field range shown in Figs. 9 and 10 are fit to Gaussian peaks. The light [dark] shaded area corresponds to $k_B T = 0.155[0.019]$ meV. Symbol styles correspond to different excitations as noted in the legend. The data points plotted with an open square with a cross inset for $\mu_0 H = 4.6$ and 5 T corresponds to the measured lower-energy excitation determined from higher energy thermally populated transitions as described in the text. Solid lines are theoretical mode energies based on finite chain calculations. Dotted and dashed grey lines in the intermediate field regime are the limiting behavior of mode energies based on finite chain calculations.

Figure 4 (a) shows the band of magnetic excitations near J_1 is also qualitatively different in the intermediate field regime. While a sinusoidal dispersion remains, the corresponding amplitude is substantially reduced (Fig. 4 (b)). This can be seen as integral to the reversion of the dispersion relation from that of magnons within a singlet for $H < H_{c1}$ where $\tilde{q} = 2n\pi$ is a minimum to magnons within a ferromagnet where $\tilde{q} = 2n\pi$ is a maximum in the dispersion relation for this mode. This change in phase of the dispersion relation is clearly seen in Fig. 13 which summarizes the dispersion relation for all fields probed in this experiment through the positions of peaks in gaussian fits to constant- \tilde{q} cuts through the data. The constant- \tilde{q} cuts were made after averaging about the $\tilde{q} = 2\pi$ wave-vector to improve counting statistics. For comparison we fit the two measurements at $T = 0.224$ K $H = 0$ data in Fig. 13(a) to the dispersion of Eq.8 and determine $J_1 = 0.44(1)$ and $\tilde{J}_2 = 0.11(1)$ meV for the coarser resolution configuration and $J_1 = 0.43(1)$ and $\tilde{J}_2 = 0.12(1)$ meV for the fine resolution configuration. We note that because a similar temperature was used

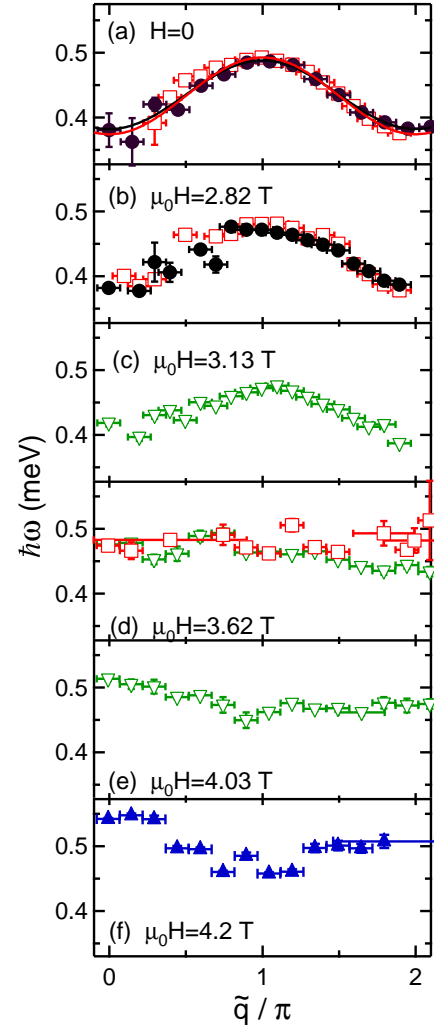


FIG. 13: (color online) Zero field (a) and intermediate field (b)-(f) dispersion relations for the $S_z = 0$ mode near $J_1 = 0.44$ meV extracted by fitting the time-of-flight measurements shown in Fig. 7(a)-(f) as described in the text. The horizontal error bar represents the range of integration in $\frac{\tilde{q}}{\pi}$ used for the constant wave-vector scans which were fit to extract the dispersion values. Multiple data sets for a given magnetic field were extracted from the different configurations used during the measurement. The blue triangles correspond to fitting the data from configuration 4. The green inverted triangles were measured in configuration 2. The solid(open) black(red) circles(squares) correspond to fits to data acquired in configuration 3(1). The solid lines in (a) are fits of the data to the zero field dispersion as described in the text.

for this measurement, the exchange values are in better agreement with the low-temperature values determined in reference 61.

More than a gradual change in the dispersion relation, Fig. 4(a), Fig. 5, and Fig. 7(c)-(f) show the character of the magnetic excitation spectrum is modified in the intermediate field range. The spectrum is broadened beyond resolution and while thermal effects surely play a role for

$T = 1.8$ K the broadening clearly visible in Fig. 7(d)-(e) for $T = 0.224$ K is an intrinsic feature of the low T spectrum.

For $H < H_{c1}$ the excitation near J_1 is associated with an $S_z = 0$ magnon propagating through the singlet ground state. The absence of damping there indicates coherent propagation through the singlet which might be considered a consequence of the lack of lower energy states for the magnon to decay into. The substantial damping for $H_{c1} < H < H_{c2}$ is evidence that the ground state has fundamentally changed character into one that offers a plethora of decay processes for the $S_z = 0$ magnon. The incoherent magnon propagation is consistent with a Luttinger liquid ground state at intermediate fields. Such a state features a continuum of low energy two-spinon excitations that can exchange linear and angular momentum with the magnon and lead to the incoherent nature of the $S_z = 0$ state that we observe.

VI. CONCLUSIONS

We have presented a comprehensive study of the wave-vector dependence of the gapped excitation spectrum of the 1d quantum spin liquid CN. A careful examination of the three dimensional dispersion of the zero field singlet-triplet excitations has established that the alternating spin chains in this material extend along the $[111]$ or $[\bar{1}\bar{1}1]$ real space direction and that the leading inter-chain interaction is $\tilde{J}'_2 = -0.01(2)$ meV.

Below the lower critical field and above the upper critical field, the excitations are well described as magnon-

like wave packets traveling through respectively a singlet ground state and a forced ferromagnet. While the phase of the \tilde{q} -dependent scattering intensity and dispersion relation is reversed the data are described by a consistent set of exchange constants. Exact diagonalization calculations for finite length alternating spin chains provide an excellent guide to understanding the field dependent data.

The intermediate field regime is considerably more complicated as it features a gapless excitation spectrum in the low T limit. Our measurements of the $\hbar\omega \approx J_1$ mode show that the change in phase of its dispersion relation is accompanied by the loss of coherence for $S_z = 0$ magnon propagation. Employing the $S_z = 0$ magnon to probe the low energy excitation spectrum, this result provides indirect evidence for a gapless continuum of excitations from the partially magnetized ground state. Given the good knowledge of the spin-hamiltonian for CN, it would be of great interest to see the numerical and analytical methods of one-dimensional quantum magnetism attempt to account for these observations.

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