Quasi-one-dimensional exchange interactions and short-range magnetic correlations in CuTeO$_4$

Zubia Hasan,1,* Eli Zoghlin,2,3 Michal Winiarski,3 Kathryn E. Arpino,4 Thomas Halloran,1 Thao T. Tran,5 and Tyrel M. McQueen1,6,†

1Institute for Quantum Matter, Department of Physics and Astronomy, The Johns Hopkins University, Baltimore, Maryland 21218, USA
2William H. Miller III Department of Physics and Astronomy, Johns Hopkins University, Baltimore, Maryland 21218, USA
3Faculty of Applied Physics and Mathematics, Gdansk University of Technology, Narutowicza 11/12, 80-233 Gdansk, Poland
4Max Planck Institute for Chemical Physics of Solids, 01187 Dresden, Germany
5Department of Chemistry, Clemson University, Clemson, South Carolina 29634, USA
6Department of Chemistry and Department of Materials Science and Engineering, The Johns Hopkins University, Baltimore, Maryland, 21218, USA

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CuTeO$_4$ has been proposed as a crystallographically distinct, yet electronic structure analog, of the superconducting cuprates. Here, we present a detailed characterization of the physical properties of CuTeO$_4$ to address this proposal. Fitting of magnetic susceptibility data indicates unexpected quasi-one-dimensional, antiferromagnetic correlations at high temperature, with a nearest-neighbor Heisenberg exchange of $J_1 = 164(5)$ K. Low-temperature heat capacity measurements reveal a sizable $T$-linear contribution of $\gamma = 9.58(8)$ mJ mol$^{-1}$ K$^{-2}$, qualitatively consistent with expectations for a $S = 1/2$, uniform, Heisenberg spin chain. Below $T \approx 40$ K, the susceptibility shows an upturn inconsistent with quasi-one-dimensional behavior. While heat capacity measurements show no signs of magnetic order down to low-temperature, the upturn in the magnetic susceptibility coincides with the emergence of a diffuse peak (centered at $|\mathbf{Q}| \approx 0.7$ Å) in the neutron diffraction data, indicative of persistent, short-range, antiferromagnetic order with a correlation length of $\xi = 10.1(9)$ Å at $T = 10$ K. The onset of nonlinearity and hysteresis in the isothermal magnetization curves suggests the presence of a small ferromagnetic component. This persistent, short-range order is understood in the context of structural modeling of the x-ray and neutron diffraction data that show the presence of a significant density of stacking faults. No evidence for substantive dopability is observed and CuTeO$_4$ appears, qualitatively, to have a larger band gap than predicted by density functional theory. We ascribe this finding to the inductive withdrawal effect from high-valence Te and suggest that superconductivity in copper tellurates is more likely to be found in compounds with a decreased reductive withdrawal effect from Te.

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I. INTRODUCTION

High-temperature superconductivity in copper oxides is one of the most exciting emergent phenomena to result from strong electron-electron correlation effects [1–3]. Significant progress has been made in the past 30 years in elucidating the key ingredients that underlie this behavior, with a layered crystallographic structure, two-dimensional electronic structure, strong Cu-$d$ and O-$p$ orbital hybridization, and proximity to an insulating, magnetically ordered state, all believed to be important.

Recent computational work identified CuTeO$_4$ as a previously unrecognized material containing these ingredients [7]. Figure 1(a) shows the reported crystal structure of CuTeO$_4$, alongside two other well-known cuprates, highlighting the layered nature. However, CuTeO$_4$ is distinct in that the Cu layer is built from distorted, highly angled, corner-sharing CuO$_4$ octahedra [Fig. 1(b)], with the Cu-O-Cu bond angle of 126.1° closer to that of herbertsmithite [8]. Thus CuTeO$_4$ gives a possible new avenue to expand the known cuprate families and provides an opportunity to test which features are most important in ultimately producing superconductivity. Despite being known for some time [9] there is scant experimental data on CuTeO$_4$, particularly in regard to its magnetic and electronic properties. This is likely due to the difficulty in preparation, which was reported to require hydrothermal conditions at 650 °C and 2000 bar for two months in order to produce a multiphase mixture containing small crystals of CuTeO$_4$ as a side product. It is clear that development of a simpler, less onerous, route to preparing this material is needed to allow for detailed characterization of its properties and evaluation of its status as a potentially novel cuprate superconductor.

Here, we report a novel method of synthesizing polycrystalline CuTeO$_4$ in mild hydrothermal conditions. The yellow color of the resulting powder suggests, qualitatively, a larger band gap than predicted by density functional theory (DFT) [7], which we ascribe to the inductive withdrawal effect of hypervalent Te$^{6+}$ [10]. Carrier doping efforts with our synthesis method were not successful and the large band gap most
FIG. 1. (a) Comparison of the layered structures of known superconductors YBa$_2$Cu$_3$O$_7$ [4] and La$_2$CuO$_4$ [5] to CuTeO$_4$. The transparent grey planes indicate the Cu layers. All atomic visualizations were created with VESTA [6] and only Cu-O bonds are shown. (b) Illustration of the two varieties of distorted CuO$_6$ octahedra in CuTeO$_4$ (Cu1 vs. Cu2) with the three unique bond distances labeled for each. The variation in coordination environment generates alternating twisted CuO$_4$ plaquettes in the Cu layers (grey transparent regions, corresponding to the two shorter Cu-O bonds in each octahedra). This strong buckling of the Cu layers is ultimately driven by shared oxygens with the TeO$_6$ octahedra. (c) Planar view of the Cu layer demonstrating how the corner-sharing connectivity of the alternating CuO$_4$ plaquettes creates a quasi-1D scenario, with the chains of plaquettes defined by the black Cu-O bonds. Note that the Cu-O-Cu pathways between different chains always involve one Cu-O bond that is not part of a CuO$_4$ plaquette.

likely rules out CuTeO$_4$ as a dopable host for high temperature superconductivity. These results suggest that future efforts at producing copper tellurate superconductors should look to minimize the inductive withdrawal effect by seeking materials with Te in a lower valence state. Nonetheless, enabled by our improved synthesis method, we present extensive characterization of the low temperature magnetic and electronic properties of CuTeO$_4$ through a combination of magnetic susceptibility, magnetization, and heat capacity, as well as x-ray and neutron scattering measurements.

At $T \gtrsim 75$ K, our magnetic susceptibility data can be fit to a one-dimensional (1D) model corresponding to a uniform, $S = 1/2$, Heisenberg chain with an antiferromagnetic nearest-neighbor exchange. Such a finding is justified by a detailed examination of the crystal structure, which reveals how the distorted CuO$_4$ octahedra form quasi-1D chains. Elastic neutron scattering reveals the emergence of magnetic diffuse scattering below $T = 60$ K. Since the AC magnetic susceptibility measurements are not consistent with a spin glass ground state we identify this diffuse scattering as due to the formation of short-range, AFM order. The observation of hysteresis in the isothermal magnetization curves, alongside the irreversibility seen in the zero-field cooled (ZFC) and field cooled (FC) DC magnetic susceptibilities ($T_{irr} = 40$ K), suggests the presence of a ferromagnetic (FM) component, likely due to a canting of the AFM order. Detailed analysis and modeling of our powder diffraction data indicates extensive faulting of the Cu layers. This has a clear impact on the magnetic properties, preventing the establishment of any long-range order down to $T = 0.1$ K, as shown by our low-temperature heat capacity data, despite the onset of short-range correlations at much higher temperatures. A relatively large $T$-linear contribution to the heat capacity is observed at low temperatures ($\gamma = 9.58(8)$ mJ mole$^{-1}$ K$^{-2}$), qualitatively consistent with the expectation for a uniform, $S = 1/2$, AFM Heisenberg chain.

II. METHODS

CuTeO$_4$ was initially synthesized by Falck et al. [9], under high pressure and temperature. Here we produce polycrystalline CuTeO$_4$ via a modified method conducted at significantly lower pressure and temperature. A 1:3 molar ratio of CuO (2 mmol) and Te(OH)$_6$ (6 mmol) was used to make the reaction mixture. This mixture was placed in a 23 mL Teflon lined stainless steel autoclave with 10 mL of water and 0.03 mL of H$_2$O$_2$ [11]. A magnetic stir bar was added to the Teflon cup and the autoclave was closed and placed on a magnetic hot plate in a sandbath. A temperature of 210 °C was maintained for seven days with the help of a thermocouple placed next to the autoclave in the sand. A difference of a few degrees from the thermocouple sensing
and the temperature within the autoclave is expected. Various attempts were made to adapt the synthesis method for carrier doping (e.g., substitution of Sb for Te) without success: no signs of dopant solubility in the parent phase were observed. We note here the observation that high-purity CuTeO₄ powder produced in this way is a bright yellow color.

The properties of the resulting polycrystalline powder of CuTeO₄ were characterized using a variety of methods. Phase purity was confirmed by powder x-ray diffraction data (PXRD) collected at room temperature using a Bruker D8 Focus diffractometer with a LynxEye detector and Cu Kα radiation (λ = 1.54 Å⁻¹). Rietveld refinements were carried out using TOPAS version 4.2 [12]. For magnetic properties measurements a small amount (~10 mg) of phase-pure CuTeO₄ powder was wrapped in a pouch of thin, polyethylene film and mounted inside a plastic straw. Care was taken to keep all mounting components clean to avoid contamination by extrinsic impurities (e.g., Fe in dust from the laboratory environment) which could affect the magnetic measurements. DC magnetic susceptibility data (χ ≡ M/H) were collected using the vibrating sample magnetometer (VSM) option of a Magnetic Properties Measurement System (MPMS, Quantum Design). AC magnetic susceptibility data were collected on the same instrument with a range of frequencies using an AC drive field of 3 Oe and a DC field of 100 Oe. Higher frequency data (f = 10 000 Hz) were collected with the same field parameters using a Physical Properties Measurement System (PPMS, Quantum Design). The heat capacity of a pressed pellet was measured using the semiadiabatic pulse method in a PPMS equipped with a dilution refrigerator down to T = 0.1 K. All data were collected on warming. The powder neutron diffraction (PND) data were obtained with the high-resolution diffractometer POWGEN at the Spallation Neutron Source at Oak Ridge National Laboratory. The sample was measured using frame 2 (center wavelength 1.5 Å⁻¹) for a total of 6 × 10⁷ counts. Stacking faults in the material were simulated using DIFFAX version 1.813 [13].

III. RESULTS

A. Crystal structure

Falck et al. previously identified the crystal structure of CuTeO₄ as belonging to the P2₁/n space group [9]. Rietveld refinement of this model to our PXRD data [Fig. 2(a)] shows clear discrepancies, especially in the predicted intensity of the (121) Bragg peak. Such discrepancies indicate that this model is an incomplete description of the structure and suggest the influence of crystallographic disorder. The (121) is an allowed reflection specifically due to the Cu atoms on the P2₁/n space group [9]. Rietveld refinements were carried out with TOPAS version 4.2 [12]. For magnetic properties measurements a small amount (~10 mg) of phase-pure CuTeO₄ powder was wrapped in a pouch of thin, polyethylene film and mounted inside a plastic straw. Care was taken to keep all mounting components clean to avoid contamination by extrinsic impurities (e.g., Fe in dust from the laboratory environment) which could affect the magnetic measurements. DC magnetic susceptibility data (χ ≡ M/H) were collected using the vibrating sample magnetometer (VSM) option of a Magnetic Properties Measurement System (MPMS, Quantum Design). AC magnetic susceptibility data were collected on the same instrument with a range of frequencies using an AC drive field of 3 Oe and a DC field of 100 Oe. Higher frequency data (f = 10 000 Hz) were collected with the same field parameters using a Physical Properties Measurement System (PPMS, Quantum Design). The heat capacity of a pressed pellet was measured using the semiadiabatic pulse method in a PPMS equipped with a dilution refrigerator down to T = 0.1 K. All data were collected on warming. The powder neutron diffraction (PND) data were obtained with the high-resolution diffractometer POWGEN at the Spallation Neutron Source at Oak Ridge National Laboratory. The sample was measured using frame 2 (center wavelength 1.5 Å⁻¹) for a total of 6 × 10⁷ counts. Stacking faults in the material were simulated using DIFFAX version 1.813 [13].

FIG. 2. (a) Rietveld refinement (red line) of the PXRD data (λ = 1.54 Å, black open circles), with difference curve below (grey line). The yellow shaded region highlights the (121) Bragg peak. (Inset) The image on the left shows a side-on view of the stacking in the ideal, P2₁/n, model. All of the Cu layers are the same leading to “A” stacking. The orange and green arrows show the interlayer connectivity of edge-sharing CuO₆ and TeO₆ octahedra, labeled by their distance along the c-axis. This distance is also represented by the difference in transparency. The O atoms which are only connected to Cu atoms have been omitted for clarity. The image on the right shows a top-down view of a single, unfaulted (“A-type”) Cu layer. (b) The model (solid green line) including stacking faults, as described in the text. The difference curve is shown only as a visual demonstration of the improvement in quality of the structural model. (Inset) Corresponding images as in (a) but with the addition of a faulted “B” layer, as described in the text. The image on the left shows how the addition of the fault modifies the interlayer connectivity. The image on the right shows a single, faulted (“B-type”) Cu layer.
of Figs. 2(a) and 2(b)]. While the two are similar, Cu atoms in the faulted layer have a slightly greater overlap with the TeO$_6$ octahedra of the underlying layer. As both original and faulted Cu layers have the same intralayer arrangement the connectivity of the Cu atoms within each layer remains the same. To test this possibility we used DIFFAX to simulate structural models with 0%-100% faulted Cu layers. Inspection of these models indicated that a probability of 70% original and 30% faulted Cu layers [Fig. 2(b)] significantly improves qualitative agreement with the experimental data in terms of the predicted peak intensities and widths. Note that this model was not arrived at via a least-squares minimization routine so the reported percentage of faulted layers is not truly quantitative. Applying this faulted model to our powder neutron scattering data (not shown) also resulted in a significant improvement in qualitative agreement relative to a Rietveld refinement with the ideal structure.

B. Magnetism

Temperature dependent, ZFC, DC magnetic susceptibility data for CuTeO$_4$ are displayed in Fig. 3(a). No clear signature of three-dimensional, long-range ordering is observed down to $T = 2$ K. Furthermore, there is no region in the susceptibility data that could be accurately fit with the Curie-Weiss model; such behavior is typical of quasi-1D systems [14–16]. A weak maximum, another characteristic of quasi-1D systems [17], is observed at $T_{\text{max}} = 100$ K, and we find that for $T \geq 75$ K, the data can be reasonably well fit by the expression:

$$\chi(T) = A\chi^\alpha\left(\frac{k_BT}{J_1}\right) + \chi_0,$$

where $A = \frac{\mu_0^2\mu_r^2/3}{J_1k_B}$ is an overall scaling factor and $\chi_0$ is a temperature independent contribution. $\chi^\alpha\left(\frac{T}{J_1}\right)$ is a fitting function derived from a high-temperature series expansion for the susceptibility of the AFM alternating exchange model:

$$\chi^\alpha(\alpha, t = k_BT/J_1) = \frac{e^{-\Delta^\alpha(\alpha,t)}}{4t}\mathcal{P}_{m(8)}(\alpha,t),$$

where $\Delta^\alpha(\alpha) = 1 - \frac{1}{2}\alpha + 2\alpha^2 + 2\alpha^3$ and $\mathcal{P}_{m(8)}(\alpha,t)$ is a modified Padé approximant, as described in Johnston et al. [18].

The AFM alternating exchange model describes a Heisenberg chain with two nearest-neighbor, AFM, intrachain exchange interactions, $J_1$ and $J_2$, with $J_2 \ll J_1$ and $\alpha = J_2/J_1$. Note that we follow the convention employed by Johnston et al. in which a positive sign is used for the AFM exchange constants. At $\alpha = 0$, the model recovers the isolated spin dimer model while $\alpha = 1$ recovers the uniform AFM Heisenberg chain model. The results from the best fit are given in Table I. Our fitting indicates that, for $T \geq 75$ K, CuTeO$_4$ behaves as system of spin chains with uniform $[\alpha = 1.000(1)]$ nearest-neighbor exchange interactions. This is suprising in the context of previous work, which considered the Cu layers as effectively 2D. We return to this point in the discussion. Based on the value of the scale factor, $A$, we extract $g = 1.4(2)$, which is significantly lower than the expected value of 2.2. However, $A$ is more strongly influenced by extrinsic factors (e.g., the error in the mass of the sample) than other

![Fig. 3. (a) ZFC magnetic susceptibility. The fit (solid red line) uses the AFM alternating exchange model described in the text; the dashed red line represents extrapolation of the fit function to lower temperature. The black arrow indicates the location of the weak maximum at $T_{\text{max}} = 100$ K. (Inset) Plot of the FC susceptibility $\chi_{\text{FC}}$ minus the ZFC susceptibility $\chi_{\text{ZFC}}$. The black arrow indicates the onset of irreversibility, $T_{\text{irr}} = 40$ K, as determined by the bifurcation of the FC and ZFC curves. All data were collected using $\mu_0H = 1$ T. (b) Isothermal magnetization curves collected at various temperatures for fields up to $\mu_0H = \pm 7$ T. (Inset) Closer view of the low-field region. Solid lines are a guide to the eye.](image-url)

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<th>Fit result</th>
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<td>$A$ (emu Oe$^{-1}$ mole$^{-1}$)</td>
<td>$4.7(2) \times 10^{-3}$</td>
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<tr>
<td>$J_1$ (K)</td>
<td>164(5)</td>
</tr>
<tr>
<td>$\alpha$</td>
<td>1.000(1)</td>
</tr>
<tr>
<td>$\chi_0$ (emu Oe$^{-1}$ mole$^{-1}$)</td>
<td>$-3.9(3) \times 10^{-4}$</td>
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fit results, meaning the systematic error is likely larger than what we have quoted, precluding any definitive interpretation. We note here that the magnitude of $\chi_0$ is somewhat larger than the value expected for the core diamagnetism based on ideal ion valences ($\approx -7 \times 10^{-5}$ emu Oe$^{-1}$ mole$^{-1}$ [19]). This is expected since the temperature-independent, diamagnetic contribution from the sample holder, which has not been removed, is comparable to the total raw signal from the sample and therefore has a strong influence on the fitted value of $\chi_0$.

Below the temperature regime where $\chi(T)$ can be well fit by this 1D model, an upturn in the susceptibility is visible near 40 K. To probe the character of this feature, field-cooled DC magnetic susceptibility data were taken and the subtraction ($\chi_{\text{FC}} - \chi_{\text{ZFC}}$) is shown in the inset of Fig. 3(a). These data show a clear onset of irreversibility at $T_{\text{irr}} = 40$ K, coincident with the upturn seen in the $\chi_{\text{ZFC}}$. Further magnetic characterization was carried out by collecting a series of isothermal magnetization curves ($M$ versus $H$), which are shown in Fig. 3(b). The magnetization curves are linear across the measured field range at higher temperatures, but begin to change slope and develop a slight curvature and small hysteresis below $T = 50$ K. A remnant magnetization of $1.3 \times 10^{-4}$ $\mu_B$/Cu$^{2+}$ is observed at 2 K.

As a complement to the DC data, AC magnetic susceptibility data were collected across six orders of magnitude in frequency (Fig. 4). The data show a small upturn below about 35 K, consistent with the significant change in slope of the magnetization curves observed below that temperature [see Fig. 3(b)]. Within the noise limitations of our measurements this feature does not appear to shift appreciably with frequency.

To further elucidate the nature of the low-temperature magnetism, time-of-flight (TOF) elastic neutron scattering was carried out at $T = 10$, 40, 60, and 120 K (Fig. 5). All Bragg peaks could be indexed using the reported crystallographic structure and no additional magnetic Bragg peaks characteristic of long-range order were observed at lower temperature. However, a broad, $T$-dependent, feature at low-$Q$ is indicated by the black arrow at the bottom of the figure. The (nuclear) peaks corresponding to CuTeO$_4$ are labeled by their $(hkl)$ value.

C. Heat capacity

Figure 7 shows heat capacity data down to $T = 2$ K. No sharp features associated with phase transitions are seen in this temperature range. Initial attempts to describe the data with a single Debye or Einstein term did not yield a satisfactory fit. However the data were well fit for $T \geq 4$ K by the sum of two Debye terms:

$$\frac{C_{\text{Debye}}}{T} = \frac{D(\Theta_{D1}, T)}{T} + \frac{D(\Theta_{D2}, T)}{T}. \quad (4)$$

![FIG. 4. AC magnetic susceptibility data collected with a range of AC drive frequencies ($f = 0.1 - 10\,000$ Hz) using constant magnetic field values.](image)

![FIG. 5. Elastic neutron scattering data at various temperatures, offset for clarity. The broad, $T$-dependent, feature at low-$Q$ is indicated by the black arrow at the bottom of the figure. The (nuclear) peaks corresponding to CuTeO$_4$ are labeled by their $(hkl)$ value.](image)
FIG. 6. High-temperature subtracted elastic neutron scattering data (filled black circles) at various temperatures. The conversion to scattering angle was performed using the center wavelength (\(\lambda = 1.5\) Å) for the frame 2 wavelength distribution at POWGEN. The red line is a fit to the data using the Warren line shape, as described in the text. (a) \(T = 10–60\), (b) 40–60, and (c) 60–120 K.

the phonon spectrum will become well-separated from the acoustic modes in energy. This can effectively create two separate subsystems leading to two terms in the heat capacity with different characteristic temperatures. If the optical modes are truly localized their contribution to the heat capacity can be captured by an Einstein term with the rest of the lattice contribution captured by a Debye term. However, if the optical modes possess significant dispersion than their contribution to the heat capacity can take a Debye-like character. Based on our fitting attempts this appears to be the case for CuTeO\(_4\).

Additional motivation for the two Debye term model is based on the presence of significant disorder—the stacking faults clearly indicated by the diffraction data—which should smear out the phonon dispersion curves in energy, pushing the optical modes further from an Einstein-like character.

The results of the two Debye term fit are shown in Table III; we note that the oscillator strengths sum to 5.86, close to the value of six atoms per formula unit for CuTeO\(_4\). This agreement, which was not obtained with other models, lends credence to the use of two Debye terms in fitting the data. Within this temperature range the expected Dulong-Petit value of 3\(R\) is not recovered. This can be explained by the high value of \(\Theta_\text{D2}\) since, within the Debye model, the plateau to the Dulong-Petit value occurs well above the Debye temperature.

TABLE III. Fit results from modeling the high temperature heat capacity data as shown in Fig. 7.

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<td>(\Theta_{\text{D1}})</td>
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</tr>
<tr>
<td>(s_{\text{D1}})</td>
<td>1.78(2)</td>
</tr>
<tr>
<td>(\Theta_{\text{D2}})</td>
<td>794(13) K</td>
</tr>
<tr>
<td>(s_{\text{D2}})</td>
<td>4.08(6)</td>
</tr>
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CuTeO₄ above \( T = 75 \, \text{K} \) is well-described by a uniform quasi-1D, AFM Heisenberg spin chain model. While the presence of extensive faulting of the Cu layers confirms the layered nature of the crystallographic structure, a detailed examination of the connectivity of the CuO₄ plaquettes clarifies how quasi-1D magnetic interactions can arise.

The Cu layers are composed of two types of distorted CuO₆ octahedra with distinct Cu-O bond length hierarchies. Figure 1(b) shows the basic unit of two corner-connected octahedra, where the relative rotation (i.e., buckling of the layer) is driven by the edge-sharing connection to neighboring TeO₆ octahedra. The CuO₄ plaquettes are defined by the shortest two Cu-O bonds in each octahedra and, due to the rotations, are only corner-connected to their neighbors along the [101] direction. The dominant superexchange interactions are then along this Cu-O-Cu pathway [Fig. 1(c), black line], mediated by the \( dx^2 - y^2 \) orbitals lying within the plane of individual CuO₄ plaquettes. This situation is similar to that of Na₂Cu₂TeO₆, where quasi-1D behavior is observed despite a layered, 2D, crystallographic structure; however, in that case, the intrachain exchange interactions were found to alternate strongly [\( \alpha = 0.10(1) \)] indicating chains of weakly interacting dimers [23]. A contrasting example is that of the monoclinic polymorph of SeCuO₄, where the dimensionality is further reduced, such that the magnetic susceptibility is best described by a model of isolated Cu tetramers [24]. Comparison of these compounds highlights how the dimensionality of magnetic phase behaviors in cuprates depends strongly on the details of the connectivity of the lattice and the resulting exchange interactions.

Our result of \( J_1 = 164(5) \, \text{K} \) is roughly half of that expected from the 126.1° Cu-O-Cu bond angle, which theory indicates should correspond to \( J_1 \approx 400 \, \text{K} \) [25]. This likely reflects the greater ionicity of the Cu-O bonds in CuTeO₄ due to the large inductive withdrawal effect of neighboring Te⁶⁺ ions [10]. The observed uniformity of the intrachain exchange is expected since, regardless of faulting, each individual Cu-O-Cu connection within a chain is the same.

The upturn and irreversibility observed in the magnetic susceptibility [Fig. 3(a)] indicate a deviation from quasi-1D behavior at \( T < 75 \, \text{K} \). One possibility is that this feature is associated with a spin glass transition. However, the lack of a clear, frequency-dependent peak in the AC magnetic susceptibility (Fig. 4) makes this unlikely. The temperature dependence of the broad peak observed in our neutron scattering data, which emerges around \( T_{\text{irr}} \), is consistent with magnetic diffuse scattering. That this additional scattering occurs only at low \( 2\theta \), with no additional temperature dependent scattering at high \( 2\theta \), further indicates this scattering is magnetic and not due to a structural transition. As our heat capacity data (Figs. 7 and 8) show no signatures consistent with long-range magnetic order we identify these features as due to the formation of short-range magnetic order which persists to low-temperature.

Based on the previous results it is clear that the disordered layering has a strong impact on the low-temperature magnetic properties of CuTeO₄. In an ideal 1D isotropic AFM Heisenberg model quantum fluctuations prevent the occurrence of long-range order at any temperature [26]. However, most real quasi-1D materials with isotropic Heisenberg exchange
do eventually order due to the presence of finite interchain exchange interactions. The temperature scale of this order is largely determined by the ratio of the inter- to intrachain interactions (see, for example, Table 4 in Ami et al. [27]).

The absence of long-range order down to temperatures significantly lower than the energy scale of the intrachain exchange interaction, despite the presence of short-range correlations at substantially higher temperatures, can be attributed to the high density of stacking faults. These stacking faults slightly modify the connectivity between Cu-chains in different layers (see green and orange arrows in the insets of Fig. 2). This in turn modifies the orbital overlap within the Cu-O-Te-O-Cu pathway perpendicular to the Cu layers, which, we propose, creates a distribution of superexchange interactions in this direction. This distribution acts as a source of frustration preventing the occurrence of long-range order across different Cu layers, and allowing only short-range order within the layers (at least within the temperature range explored here).

Another possible interpretation is that the distribution of exchange interactions actually induces short-range magnetic order into the system, which, in the absence of disorder, would show none. We are unable to definitively distinguish between these two interpretations due to the fact that the stacking faults are extended over an entire layer. This means that, while the disordered layering may not be the thermodynamically stable state, post-synthetic measures, such as annealing, are dynamically insufficient to resolve the structure to the $P2_1/n$ "ideal." However, this interpretation seems unlikely in the context of the many other cuprates spin chains which are observed to order (e.g., SrCuO$_2$ [28], BaCu$_2$Si$_2$O$_7$ [29], KCuF$_3$ [30], and Cu(Cu$_2$H$_2$N$_2$)(NO$_3$)$_2$ [31]). Theoretical work by Yasuda et al. [32] provides an empirical relation (equation 8 in their paper) for estimating the expected interchain coupling ($J'$) for Heisenberg spin chains based on the observed transition temperature ($T_N$) and $J_1$. Taking $T_N = 0.1$ K (the lowest temperature for which we have data) as an upper bound, this relation gives $J'/J_1 = 6.1(2) \times 10^{-4}$. This value would suggest that CuTeO$_4$ possesses some of the most isolated chains among quasi-1D cuprates [27,31], which appears unrealistic given the presence of clear Cu-O-Cu interchain exchange pathways within the Cu layers.

This distinction aside, our physical picture is the formation of small, AFM, domains of coherently ordered chains within a Cu layer (onsetting near $T_{irr}$) but with minimal coherence between distinct layers of chains. The correlation lengths from the Warren lineshape analysis at $T = 10$ K and 40 K are on the order of the interlayer spacing [5.163(4) Å], further reinforcing this picture.

Our neutron data are insufficient to definitively characterize the symmetry of this short-range magnetic order. However, the location of the diffuse magnetic scattering ($|\vec{Q}| \approx 0.7$ Å$^{-1}$) is close to the value for the (010) reflection ($|\vec{Q}| = 0.61$ Å$^{-1}$). This would indicate AFM order along the $b$ axis, perpendicular to the Cu layers, with at least some component of the moment in the (011) plane. An alternative interpretation that we cannot definitively rule out is that this scattering reflects the appearance of magnetic order with a propagation vector of $\vec{k} = (0.5, 0, 0)$, with $|\vec{Q}| = 0.57$ Å$^{-1}$ for the first peak. However, this seems unlikely as the standard crystallographic cell of CuTeO$_4$ can accommodate in-plane magnetic order without enlarging the unit cell. While the precise symmetry of the short-range order remains an open question, the finite hysteresis in the magnetization curves below $T_{irr}$ [Fig. 3(b)] indicates a weak FM component. This probably stems from a canting of the overall AFM order due to a finite Dzyaloshinskii-Moriya interaction, as has been seen in other layered cuprates [33,34].

Another relevant point is that, due to the lack of energy discrimination in our diffraction measurement, we are unable to determine if there is an inelastic component to the observed diffuse scattering; future neutron spectroscopy measurements would be of interest to address this. We note that there is a slight feature in the magnetic susceptibility near $\approx 20$ K. The origin of this feature cannot be rigorously determined with the available data. However, we believe it is most likely due to the presence of a weak paramagnetic signal—from either a small, unresolved impurity phase or defect spins within the CuTeO$_4$ matrix—which is overlaid with the saturating signal from the weak ferromagnet component we have proposed.

The linear term extracted from the low-temperature heat capacity data is surprising given that CuTeO$_4$ appears to be a large band gap insulator, precluding the possibility of a high density of electronic states at the Fermi level. However, the uniform $S = 1/2$, AFM Heisenberg spin chain is known to possess a linear heat capacity at low temperature due to gapless spin excitations [17,18,35] with the value of $\gamma$ given as $\gamma = \frac{2\pi k_b}{3J_1}$ [18]. Inputting our $J_1$ value yields $\gamma = 33(1)$ mJ mole$^{-1}$ K$^{-2}$, which is significantly higher than we observe [$\gamma = 9.58(8)$ mJ mole$^{-1}$ K$^{-2}$]. This can be explained by the reduction of $J_1$ due to the increased ionicity of the Cu-O bond, as mentioned previously. It is also possible that the influence of interchain interactions, which perturb the system towards a higher dimensionality where no $T$-linear heat capacity term is expected, may also play a role. While our low-temperature heat capacity measurements allow us to qualitatively describe the magnetic contribution to the heat capacity, a quantitative determination is not currently possible due to the lack of a nonmagnetic analog for CuTeO$_4$. However, at low temperatures, the phonon contribution to the heat capacity often scales as the dimensionality of the material. The finite $T^3$ contribution ($\beta$ term) observed here therefore reflects the relatively 3D character of the phonon dispersion.

V. CONCLUSION

While CuTeO$_4$ was predicted to have a 2D AFM ground state [7], extensive characterization enabled by our improved synthesis method demonstrates AFM Heisenberg spin chain behavior for $T > 75$ K with a uniform nearest-neighbor exchange of $J_1 = 164(5)$ K. Careful consideration of the Cu connectivity indicates this behavior is due to the arrays of corner connected CuO$_4$ plaquettes highlighted in Figs. 1(b) and 1(c). Although no long-range order was detected in our heat capacity measurements, the combination of an upturn in the magnetic susceptibility and diffuse scattering in the neutron diffraction data indicate persistent short-range magnetic order below $T = 60$ K. The persistence of these short-range
correlations down to temperatures well below the energy scale of the intrachain exchange interaction is attributed to the presence of a significant fraction of stacking faults. These faults modify the interlayer connectivity and create a distribution of interchain interactions that frustrates any incipient magnetic order. The low-temperature heat capacity shows an unexpectedly large linear term for a wide band gap insulator which is likely related to the quasi-1D magnetism.

Although the first principles calculations in Botana et al. [7] showed CuTeO$_4$ to have a small bandgap of 0.13 eV, making it a suitable candidate for superconductivity upon doping, this appears inconsistent with the yellow color of the polycrystalline powder. The underestimation of the band gap may be due to the failure of DFT to consider the effect of hypervalent cations like Te$^{6+}$. Additionally, we find no evidence of dopant solubility with our synthesis method. These findings suggest that future efforts towards discovering high-temperature superconductivity in copper tellurates should be focused on cases where there is a reduced inductive withdrawal effect by Te, such as those compounds where Te is present in the 4$^+$ state.

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[11] The inclusion of H$_2$O$_2$ seems to be crucial for the formation of CuTeO$_4$ in such mild conditions. Repetition of this experiment without H$_2$O$_2$ resulted in the formation of Cu$_3$TeO$_6$, the more thermodynamically favorable phase. The decomposition of CuTeO$_4$ into Cu$_3$TeO$_6$ alongside Te$_2$O$_3$ and O$_2$ at 510 °C is documented in [36]. H$_2$O$_2$ quenches this decomposition by preventing the formation of Te$_2$O$_3$, oxidizing Te$^{4+}$ to Te$^{6+}$.


