Quantum Disordered Ground State in the Heisenberg-Kitaev Candidate NaRuO2

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Quantum Disordered Ground State in the Heisenberg-Kitaev Candidate $NaRuO_2$

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The realization of spin liquid states born from the near-critical regime of the triangular lattice Hubbard model in inorganic materials remains a long-standing challenge, where weak spin-orbit coupling and other small perturbations often induce conventional spin freezing or order. Strong enough spin-orbit coupling, however, can renormalize the electronic wave function and induced anisotropic exchange interactions that promote magnetic frustration. Through the cooperative interplay of spin-orbit coupling and correlation effects, here we show that the triangular lattice magnet $NaRuO_2$ hosts an inherently fluctuating magnetic ground state with thermodynamic properties suggestive of a crossover between dynamic ground states. Despite the presence of a charge gap, we find that low-temperature spin excitations generate a metal-like term in the specific heat and continuum excitations in neutron scattering, reminiscent of spin liquid states found in triangular lattice organic magnets. Further cooling reveals that these fluctuations crossover into a state whose dynamic spin autocorrelation function reflects persistent fluctuations within a highly disordered spin state. These findings instantiate $NaRuO_2$ as a unique, Heisenberg-Kitaev cousin to organic, Heisenberg spin liquid compounds with a low-temperature crossover in quantum disorder driven via the interplay between geometric frustration, extended hopping, and relativistic spin-orbit coupling.

The interplay between electron-electron correlation effects and geometrical frustration can lead to a rich hierarchy of electronic states. The Hubbard model at half-filling shows that, as the on-site Coulomb interaction (U) is reduced relative to the electron hopping energy (t), the ground state transitions from an antiferromagnetic insulating state into a nonmagnetic metal. If the underlying lattice is triangular, then an intermediate nonmagnetic insulating state is predicted prior to the onset of the metallic state[1,2,3]. The predicted properties of this intermediate phase vary depending on the theoretical approach, however it is generally envisioned as an inherently quantum disordered magnetic state or a quantum spin liquid phase generically realized at the boundary of the Mott insulating regime.

Proposed physical manifestations of this type of spin liquid state are rare, with well-studied candidates identified in anisotropic triangular magnets built from organic molecular complexes[4,5,6,7]. In inorganic compounds, however, triangular lattice systems identified thus far are located either deep in the insulating regime,[8] deep in the metallic state,[9] or better described in the strong coupling limit via a pure

Heisenberg model.[10] One promising means of more fully exploring this materials space is to consider compounds with extended *d*-electron orbitals, where the cooperative interplay between moderate on-site Coulomb repulsion (U) and spin-orbit coupling (λ) driven bandwidth narrowing can generate marginally stable J_{eff}=1/2 Mott insulating states with $U/t \approx 1$.[11] Such states are known to form in *5d*-transition metal iridates, where controlling the bandwidth can drive a metal-insulator transition,[12] and similarly, weak J_{eff}=1/2 Mott states may also be realized in *4d*-transition metal compounds.[13]

Specifically, Ru^{3+} ($4d^5$) ions in a nearly cubic crystal field are capable of assuming a low-spin state with both λ and U appreciable enough to stabilize a half-filled J_{eff}=1/2 orbital.[14,15] An ideal triangular lattice decorated by octahedrally coordinated Ru^{3+} moments is also known to form in delafossite variants such as $NaRuO_2$,[16] which suggests an opportune setting for searching for an intermediate quantum disordered state at the boundary of the J_{eff}=1/2 Mott state's stability. Remarkably little is known regarding the ground state of this compound and, more generally, whether it can provide a suitable experimental window into the electronic properties of a near-critical triangular lattice Hubbard model in the presence of strong λ.

Here we establish that $NaRuO_2$ hosts electrons in a unique interaction space as an ideal triangular lattice possessing a weak spin-orbit assisted Mott insulating ground state—one consistent with the expectation of a near-critical J_{eff}=1/2 Mott state driven by cooperative λ and U. Despite the presence of a charge gap, the high-temperature magnetism of $NaRuO_2$ shows an elevated Pauli-like susceptibility, and at low temperatures, where the charge gap is well established, heat capacity data show a substantial linear term akin to the Sommerfeld coefficients expected in metals. The magnetic excitations comprising this linear term generate a diffuse continuum of spin excitations, a portion of which are slowed down below a field-coupled crossover into a state with persistent fluctuations and a heat capacity quadratic in temperature. Our data demonstrate that J_{eff}=1/2 electrons built from extended *d*-orbitals on a triangular lattice stabilize quantum disordered magnetic phase behavior, consistent with expectations of Hubbard models near the metal-insulator phase boundary[17] as well as models of Kitaev antiferromagnets.[18,19]

Figure 1 (a) shows the detailed crystal structure of $NaRuO_2$. Edge-sharing RuO_6 octahedra form a triangular lattice of Ru^{3+} ions separated by planes of Na ions. An ideal equilateral triangular lattice results and a slight (< 5°) trigonal distortion exists in the oxygen octahedra surrounding the Ru ions. Neutron diffraction data presented in Fig. 1 (b) show fully occupied Na sites and no site mixing with the resolution of the measurement. Synchrotron powder diffraction measurements (ESI) corroborate these conclusions as well. Furthermore, neutron powder diffraction data collected at temperatures as low as 50 mK show no signs of structural symmetry breaking or static spin correlations forming between Ru moments. This is remarkable given the large covalency and enhanced exchange expected between the extended Ru moments, which are only 3.06 Å apart.

Ab initio electronic structure calculations shown in Fig. 1 (c) reveal that the band structure of $NaRuO_2$ is highly sensitive to the structural parameters. Using the experimentally-derived structure in this paper, a charge gap forms via inclusion of both λ and U using the local density approximation (LDA). Choosing U=1 eV, consistent with prior LDA + λ + U models of Ru^{3+} systems such as $RuCl_3$,[20] opens a gap when incorporated with λ. However, the model and the prediction of the presence or absence of a gap is highly sensitive to the local distortions of oxygen octahedra around the Ru sites—consistent with the notion of a marginally insulating state where extended hopping effects can play an important role. The resistivity data presented in Fig. 2 (a) verify that $NaRuO_2$ indeed possesses an insulating ground state whose electrical transport is best modeled via a two-dimensional variable range hopping; however we note that

discrimination between two-dimensional and three-dimensional forms is difficult in polycrystalline samples.

Contrasting the electrical transport data, the temperature-dependent magnetic susceptibility data presented in Fig. 2 (b) instead show a nearly itinerant magnetic response. The weak increase in the susceptibility upon cooling does not fit to a conventional Curie-Weiss form, and instead, the low-temperature susceptibility is best fit to a large Pauli-like term (χ_0) with a weak concentration of free impurity moments (1.8±0.2% of S=1/2 moments). Notably, the χ_0 term is unusually large for an insulator, χ_0=9.1±0.2×10^{-4} emu Oe^{-1} mol^{-1}, a value comparable to Pd metal.[21] The large χ_0 fit via the temperature dependent magnetization matches the linear term found in the isothermal magnetization (χ_0=1.2±0.1×10^{-3} emu Oe^{-1} mol^{-1}), which can be well parameterized by a dominant Pauli-like linear term and a small fraction of S=1/2 moments (1.4±0.2%) shown in Fig. 2 (c). At lower temperatures, a weak cusp appears in the susceptibility data below 1.5 K, as shown in the inset of Fig. 2 (b). There is a weak frequency dependence to the onset of this cusp as shown in Supplemental Figure 2, which deviates from the expectations of a canonical spin glass, but nevertheless suggests a weak freezing in the local moment contribution to the overall susceptibility. The nature of this slowdown in the dynamics will be discussed later in this paper.

Exploring this further, heat capacity data were collected down to 80 mK and are plotted in Fig. 2 (d). As the system is cooled far below the Debye temperature, $C_p(T)$ takes on the form $C_p(T)$=Aγ+βT^3 where β parameterizes the contribution from phonons and γ is an anomalous linear term typically observed in metals. In free electron gases, γ represents the Sommerfeld coefficient—a term proportional to the electronic density of states occupied at the Fermi level; however in magnetically frustrated insulators the presence of a γ term suggests a fractionalization of electrons and the presence of a spinon Fermi surface. The γ term in $NaRuO_2$ is substantial (12.5±0.07 mJ mole^{-1} K^{-2}) and is distinct from the small γ's found in disordered insulators. Instead, it is comparable in magnitude to those found in organic triangular lattice spin liquid candidates.[22,23,24] One notable distinction is that the presence of strong spin-orbit coupling amplifies χ_0 and the resulting Wilson ratio $R_W = \frac{\pi^2 k_B^2 \chi_0}{3\mu_B^2 \gamma}$ = 8 for $NaRuO_2$ versus $R_W \approx 1$ for organic compounds.

Cooling below 2 K reveals the onset of a weak freezing transition in $C_p(T)$ matching the cusp in the low-T susceptibility. Only a small amount of entropy is associated with this T_F = 2 K crossover (1.2±0.2% of Rln(2)) further correlating it with a freezing of a small fraction of local moments. However, T_F also marks the onset of a modified power law $C_p(T)=AT^{\alpha}$ that depends on the magnetic field. Applying a magnetic field shifts this T_F crossover upward in temperature, and, as the shoulder of this freezing feature shifts to higher temperatures, the low temperature $C_p(T)$ converges to a quadratic behavior (α = 2). Notably, the application of a magnetic field also enhances the apparent γ term above T_F, and it increases by 18% under $\mu_0 H$ = 14 T.

To investigate the nature of the magnetic excitations associated with these low energy fluctuations, inelastic neutron scattering measurements were performed. As an initial survey, temperature subtracted data (2 K -300 K) are plotted in Fig. 3 (a) where magnetic spectral weight at low momentum transfers appears centered near 25 meV. Energy cuts (integrated over small momentum transfers Q) are plotted for both the 2 K and 300 K datasets in Fig. 3 (c) and show a peak in scattering at 2 K that is broadened, strongly diminished, and shifted downward in energy upon warming to 300 K. Direct subtraction of the

type shown in Fig. 3 (a) often oversubtracts the low-energy portion of magnetic scattering due to the thermal population of low energy phonons. To compensate for this, a Bose-corrected subtraction of the data for a lower incident energy is also shown in Fig. 3 (b). The momentum distribution of the resulting low-energy continuum is centered near 1.3 $Å^{-1}$ as shown in Fig. 3 (d)

To probe the spin dynamics at even lower frequencies and temperatures, the muon spin relaxation technique was used for the determination of different static local fields and the presence of magnetic fluctuations. The temperature dependent muon polarization and their fit are presented in Fig. 4(a). At 12 K, the muon spin depolarization is best described via a Gaussian Kubo-Toyabe form dominated by the contribution of nuclear moments in the sample. Upon cooling below 12 K, the magnetic fluctuations from the antiferromagnetically coupled electronic spins slow down into the muon time window and the time-dependent polarization is described by a generalized depolarization function $P(t) = (f)GbG(\Delta; R; t)e^{(-\lambda_{GbG}t)^{\beta}} + (1-f)e^{-\lambda_p t}$. This is a response comprised of an uncorrelated fraction $(1-f)$ of paramagnetic moments whose fluctuations drive simple exponential relaxation at λ_p that, upon cooling, converts into a highly disordered response captured by a Gaussian-Broadened-Gaussian (GbG) function dressed by spin fluctuations in the form of a stretched exponential with relaxation rate λ_{GbG}. The GbG function[25] represents a normal distribution of Gaussian field distributions about a central value Δ_0 with width w captured by the parameter $R = w/\Delta_0$. As shown in Fig. 4(b), Δ_0 increases progressively with decreasing the temperature below 3 K, and R peaks at approximately 2.5 K, with the value changing from 0 at high temperatures to 1 at the base temperature. The absence of clear oscillatory signals suggests that the low-temperature state contains intermediate-diluted and disordered static magnetic moments. Meanwhile, we notice that this nominally static distribution of fields is dressed by a slow fluctuation rate (λ_{GbG}). As presented in Fig. 4(c), similar to other highly frustrated materials with persistent spin fluctuations in their ground states[26]. To discriminate the presence of persistent fluctuations in this disordered ground state versus depolarization via static disorder, a longitudinal field (LF) experiment was performed. Such a field rapidly decouples muons from slow depolarization due to purely static field distributions, whereas depolarization persists in systems that are inherently dynamic[27]. The data in Fig. 4 (d) instead show the persistence of fluctuations under modest applied fields, for example, contrasting the rapid decoupling seen in static field distributions in frustrated, yet frozen systems such as $Na_4Ir_3O_8$.[28,29]

To further distinguish the partial freezing and persistent dynamics in $NaRuO_2$ from conventional spin glass behavior, we demonstrate the magnetic-field time scaling relations of the LF time-spectra using the form $P(H, t)=P(t/H^{\gamma})$.[30] The polarization maps to the spin autocorrelation function, and, above the freezing temperature, the depolarization should scale as a function of t/H^{γ} with the scaling exponent γ reflective of the manner through which spin correlations decay in time.[31] With the exception of the short-time limit μ_0H = 0.02 T data, the data scale well to this form with the analysis shown in Figs. 4 (e) and (f). This scaling analysis yields an exponent of $\gamma \approx 1.75$, which precludes conventional spin-glass power law correlations ($\gamma \leq 1$) and, instead, is consistent with the expectation of decay via a stretched exponential function—identical to the type used to model the zero-field data.

The above experimental picture suggests that $NaRuO_2$ occupies a unique phase space where a spin-orbit assisted Mott state gives rise to a native quantum disordered ground state. Anisotropic Kitaev interactions are expected to play an important role in this disordered state. Ru^{3+} ions sit in a nearly cubic crystal field—one that should promote the formation of a J_{eff}=1/2 wavefunction from the t_{2g} orbital

manifold. An array of such wavefunctions on a honeycomb lattice composed of edge-sharing octahedra generates strong Kitaev exchange and is predicted to stabilize a quantum spin liquid ground state, but experimental realizations are lacking. Although materials such as Na_2IrO_3[32] and $RuCl_3$[33] are known to stabilize strong ferromagnetic Kitaev coupling strengths, due to dominant Heisenberg interactions they nevertheless possess magnetically ordered ground states.[34,35] Similar predictions of dominant Kitaev exchange and quantum spin liquid phases have also been put forward for J_{eff}=1/2 electrons decorating a triangular lattice.[19] Unlike other Kitaev candidate materials, the thermodynamic phenomenology of $NaRuO_2$ resembles that of highly frustrated organic spin liquids. We hypothesize this is due to the weak nature of the insulating state and consequent importance of longer-range exchange interactions. These ingredients are believed to stabilize intrinsically quantum disordered ground states[36,37] as we propose is realized here.

Inorganic compounds with simple structures and large magnetic exchange energy scales rarely realize quantum disordered or spin liquid-type ground states due to low-energy interactions, such as Dzyaloshinskii-Moriya interactions or weak lattice distortions that lift magnetic frustration. Even rarer is a quantum spin liquid candidate that manifests a transition/crossover between quantum disordered phases manifest in the breakdown of the γ term in $NaRuO_2$. The unique interplay between the marginally insulating Mott state and anisotropic exchange fostered by cooperative spin-orbit coupling and on-site Coulomb interactions suggests that $NaRuO_2$ occupies a unique energy landscape—one where Kitaev coupling in the Hubbard model fosters nearby quantum disordered states. The result is a material that provides an appealing opportunity for pushing the manifestation of exotic phenomena such Majorana excitations associated with a native Kitaev spin liquid state up to higher temperatures for future quantum information applications.

Methods:

Sample preparation: Sample preparation: Polycrystalline $NaRuO_2$ was synthesized using mechanochemical methods. Sodium peroxide (Na_2O_2) beads (Sigma, 97%), ruthenium dioxide (RuO_2) powder (Alfa, 99.95%), and Na metal (Alfa 99.8%) are combined in a pre-seasoned tungsten carbide ball-mill vial under argon. The synthesis generally needs excess Na and O during the reaction; though we note that $NaRuO_2$ exhibits an unusually large degree of off-stoichiometry in the Na-rich regime. To optimize purity and Na-stoichiometry, we have designed the synthesis to minimize Na_{Ru} antisite defects by forcing $NaRuO_2$ to crystallize in equilibrium with Ru and $NaRu_2O_4$, which corresponds to the Ru-rich edge of the phase diagram. Thus, the stoichiometry has been tuned empirically to $Na_{1.07}(Na_2O_2)_{0.35}(RuO_2)_{1.35}$. A more detailed analysis of the phase diagram and the resulting Na_{Ru} off-stoichiometry will be published elsewhere.

The resulting mixture is milled for 60 minutes in a Spex 8000D Mixer/Mill using four 7.9mm tungsten carbide balls. The reaction generates a substantial amount of heat, though powders produced through milling are effectively amorphous. Resulting powder is lightly ground in an agate mortar, sieved through a 100 micron sieve, and loaded into 2mL CoorsTek alumina crucibles. The crucibles are sealed under approximately 1 bar of argon in fused silica ampoules and immediately placed within a preheated furnace at 900°C. Samples are annealed for 30min and are then immediately air-quenched before extracting

powders within an argon glovebox with water and O_2 levels <0.5ppm. Resulting powders are largely phase pure with trace amounts of Ru metal (<2%). Powders are black and are highly moisture sensitive.

X-ray Synchrotron diffraction: High-resolution synchrotron powder diffraction data were collected using beamline 11-BM at the Advanced Photon Source (APS), Argonne National Laboratory using an average wavelength of 0.457925 Å. Both 300 K and 5K measurements were performed to check for any crystallographic phase transformations, and for better analysis of thermal parameters and occupancies at base temperatures. Discrete detectors covering an angular range from -6 to 16° in 2θ are scanned over a 34° range, with data points collected every 0.001°, and scan speed of 0.01°/s. Due to the air sensitivity of the materials, small quantities of $NaRuO_2$ were diluted with amorphous SiO_2 in a glovebox and sealed under argon in flame-tipped amorphous SiO_2 capillaries. These capillaries were nested within kapton sleeves and held in place with a small amount of modeling clay. Diagnostic laboratory powder diffraction utilized a Panalytical Empyrean diffractometer (Cu Kα, 1.54 Å) in the Bragg-Brentano geometry.

Neutron diffraction: Neutron powder diffraction measurements were performed on the fixed incident energy triple-axis spectrometer HB-1A (λ=2.37 Å) of the High Flux Isotope Reactor (HFIR) at ORNL. 6 grams of phase pure polycrystalline $NaRuO_2$ was loaded in a cylindrical Cu can which was then placed in a dilution insert of an orange cryostat, providing a base temperature of 50 mK. The collimation configuration of 40'-40'-40'-80' selected yielded an energy resolution (FWHM) at the elastic line of ~ 1 meV. The combination of the double-bounce monochromator system and the placement of the pyrolytic graphite (PG) crystal analyzer for energy discrimination before the single He-3 detector provided an excellent signal-to-noise ratio. Additional contamination from higher-order wavelength contamination was minimized with the use of two PG filters.

High resolution neutron powder diffraction experiments were performed on the high resolution neutron powder diffractometer BT-1 of the NIST Center for Neutron Research (NCNR) at the National Institute of Standards and Technology (NIST). A Ge(311) monochromator provided a λ=2.0772 Å with maximum neutron flux, allowing for full diffraction patterns to be collected in 12 hours. 4.5 grams of phase pure polycrystalline $NaRuO_2$ was loaded into a vanadium can under a He atmosphere and placed into a top loading flow-type orange cryostat, providing a base temperature of 1.6 K. Rietveld refinements of the neutron diffractograms were performed with TOPAS Academic V6.

Inelastic neutron scattering: High-energy inelastic neutron scattering experiments were performed on the direct geometry time-of-flight chopper spectrometer SEQUOIA of the Spallation Neutron Source (SNS) at Oak Ridge National Laboratory.[38] 8 grams of phase pure polycrystalline NaRuO2 was loaded in a cylindrical Al can which was placed in a top loading helium cryostat, providing a base temperature of 1.8 K. Powder-averaged (Q,E) spectra were collected with incident energies of 80 meV and 43 meV, operating in high flux mode, providing an elastic resolution of ≈7.8% and ≈2.3% E_i, respectively. Background contributions to the inelastic spectra were approximated by an empty aluminum, which was measured in identical experimental conditions for approximately one third of the counting time allocated for the sample.

Magnetic Susceptibility (MPMS, PPMS, and ACDR): The temperature dependence of the zero-field-cooled (ZFC) and field-cooled (FC) DC magnetization of 17.6 mg of phase pure $NaRuO_2$ powder placed in a polystyrene holder was measured on a 7 T Quantum Design Magnetic Property Measurement System (MPMS3) SQUID magnetometer. Data was collected continuously in sweep mode with a ramp rate of 2 K/min in the presence of an external DC field of 1000 Oe (1 Oe = (1000/4π) A/m).

Isothermal magnetization measurements of the same 17.6 mg $NaRuO_2$ sample at 1.8 K and 300 K were performed on a Quantum Design 14 T Dynacool Physical Property Measurement System (PPMS) employing the vibrating sample magnetometer (VSM) option. ZFC data was collected continuously in sweep mode with a ramp rate of 100 Oe/sec .

The temperature dependence of the ac magnetization of 10.1 mg of phase pure $NaRuO_2$ powder was measured on a Quantum Design 14 T Dynacool PPMS employing the ac susceptibility option for the dilution refrigerator (ACDR). The milled powder was cold pressed with a Carver press, heated at 900 ^oC, and a portion of the sintered pellet with approximate dimensions of 1 mm x 1 mm x 0.5 mm was adhered to a quartz sample mounting post with a thin layer of GE-varnish. All ac measurements were collected in stable mode under ZFC conditions in an external DC magnetic field of 1 T and with a ramp rate of 0.08 K/min.

Heat Capacity: The temperature and field dependence of the specific heat capacity of 5.62 mg of phase pure $NaRuO_2$ powder and 6.21 mg of its non-magnetic analog $NaRhO_2$ were measured on a Quantum Design 14 T Dynacool PPMS employing the heat capacity (HC) option. Both $NaRuO_2$ and $NaRhO_2$ powders were first cold pressed with a Carver press, heated at 900^o C, and a portion of the sintered pellets of approximate dimensions 0.5 mm x 0.75 mm x 0.35 mm was adhered to the sample stage with Apiezon N grease to optimize thermal coupling. All measurements were performed upon heating, while all measurements in field were done in ZFC conditions.

SPS: $NaRuO_2$ pellets (10 mm diameter x 2 mm thick) were prepared from phase pure $NaRuO_2$ powder using field-assisted sintering (FCT Systeme GmbH, Frankenblick, Germany). The pellets were pressed at 850 °C and 90 MPa for 60 min in an Ar-filled chamber with a pressure of 30 hPa, using a heating rate of 150 °C/min and a cooling rate of 40 °C/min. All pellets were subsequently ground to a 2000-grit finish before resistivity measurements.

Resistivity: $NaRuO_2$ pellets were then sectioned into rectangular bars with dimensions approximately 1 mm x 2 mm x 0.5 mm. Electrical contacts were made in a standard four-point geometry with contacts being made with a combination of gold wire and silver paint. Thermal contact and electrical isolation was ensured using layers of GE varnish and cigarette paper. The temperature dependence of the electrical resistivity was measured with the Electrical Transport Option (ETO) on a 9 T Quantum Design Dynacool PPMS using a drive current of 10 μA and drive frequency of 100 Hz. Data was collected continuously in sweep mode with a ramp rate of 2 K/min.

Muon spin relaxation measurements: Muon depolarization data were taken on the General Purpose Surface-muon spectrometer on the πM3.2 beamline at the Paul Scherrer Institute. For the zero field data the muon beam spin polarization was oriented at 45° to the muon momentum, while for longitudinal field studies the polarization was anti-parallel to the momentum. A pressed powder disk of diameter 10 mm and thickness 2 mm was wrapped in thin mylar foil and suspended in a gas-flow cryostat. Data were fit using the MUSRFIT program[39].

Electronic structure calculations: First-principles electronic calculations based on density functional theory (DFT) were performed using the Vienna ab initio Simulation Package (VASP version 5.4.4) [40,41] with the local density approximation functional and projector-augmented wave (PAW) pseudopotentials.[42,43] The plane-wave energy cutoff was set to 400 eV and a Γ-centered 15 × 15 × 15 k-point mesh was automatically generated within VASP. Initial biasing of the spin-polarization and tetrahedral smearing with

Blöchl corrections were used for the self-consistent static calculation.[44] The electronic structure was calculated via a non-self-consistent run using the charge density from the static calculation with spin-orbit coupling (SOC) and a Hubbard U correction of 1 eV. A k-point path for the band structure calculation was generated using the AFLOW online tool.[45] All calculations had energy convergence better than 10^{-6} eV.

Data Availability: The data that support the findings of this study are available from the corresponding author upon reasonable request.

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Author Contributions: S.D.W. wrote the manuscript with input from all coauthors. B.R.O., P.S., and A.H. synthesized the material and performed resistivity, magnetization, and neutron scattering measurements. A.H. and R.S. performed *ab initio* DFT calculations. E.K., M.J.G., and C.W. performed muon spin relaxation measurements. A.K., C.M.B., and A.A. performed neutron scattering measurements. L.B. provided theoretical insight into modeling the material. All authors participated in planning and discussions of experiments.

Additional Information: Supplementary Information accompanies this paper.

Competing interests: The authors declare no competing financial or non-financial interests.

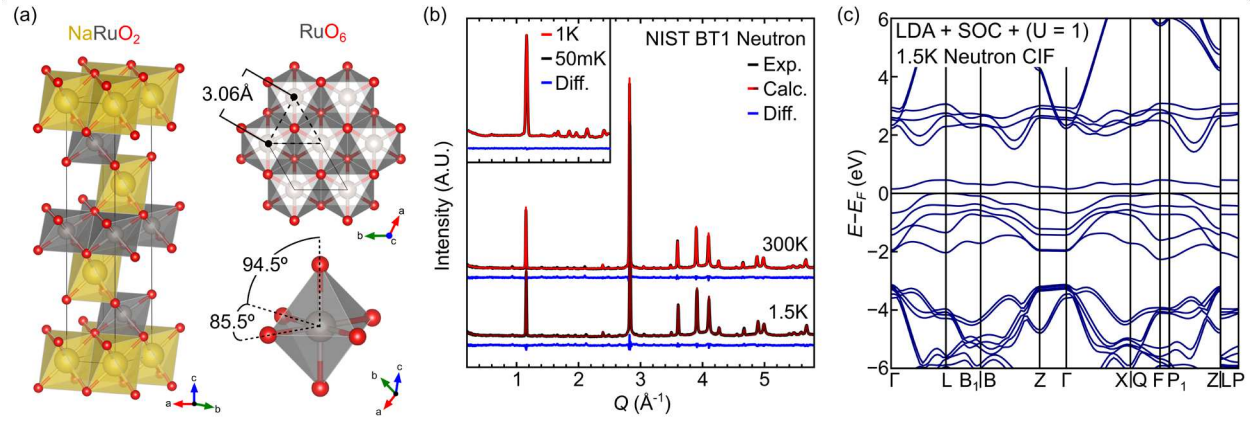

Figure 1: Lattice and electronic band structures of $NaRuO_2$ (a) Crystal lattice of $NaRuO_2$ showing the triangular lattice of edge-sharing RuO_6 octahedra arranged in a layered structure separated by planes of Na-ions. The slight trigonal distortion of the RuO_6 octahedra is illustrated. (b) Neutron powder diffraction data collected at 300 K, 1.5 K, 1 K, and 50 mK. Data in the main panel were collected at BT-1 and are overplotted with the refined structural diffraction pattern. The inset shows data collected on HB-1A and reveals no magnetic scattering appearing down to 50 mK. (c) LDA electronic band structure for $NaRuO_2$ using both λ and U=1 eV.

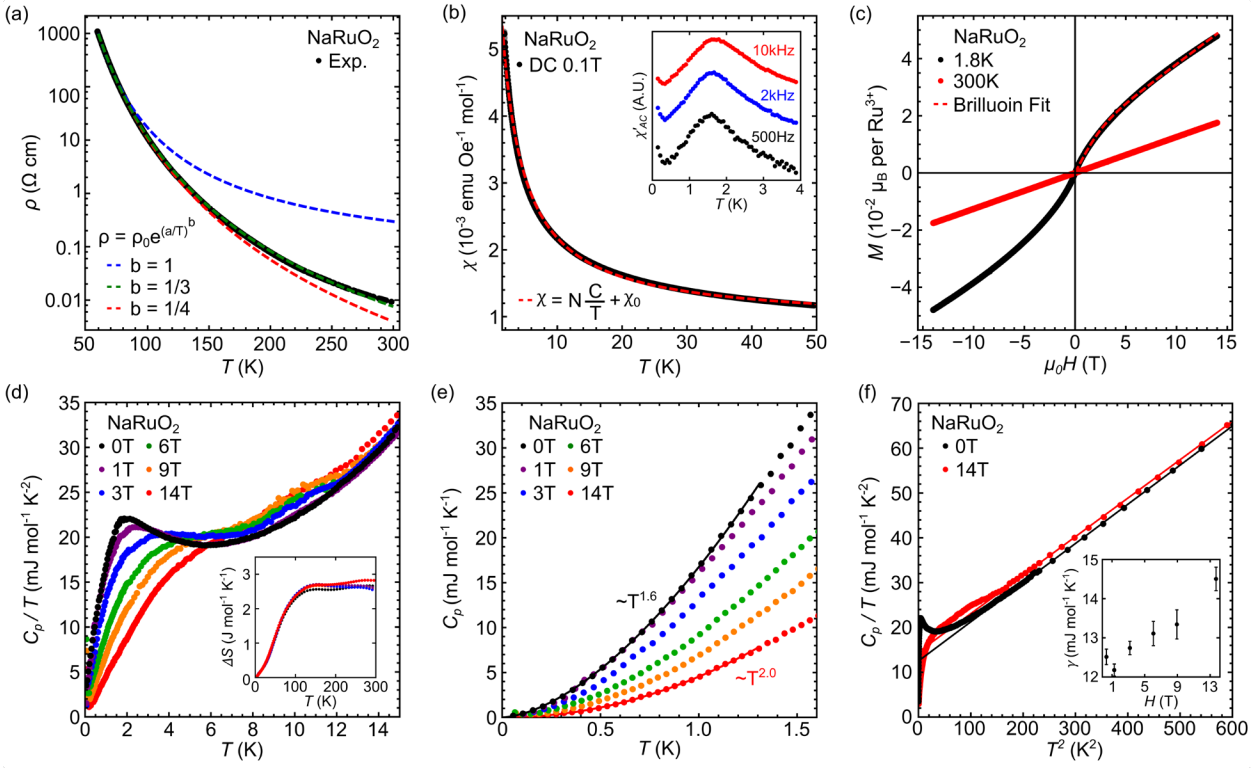

Figure 2: Electrical transport, magnetic susceptibility, and heat capacity data characterizing the low temperature properties of $NaRuO_2$ (a) DC resistivity data collected as a function of temperature. Fits to models of variable-range hopping transport are overpotted for both two-dimensional (b=1/3) and three-dimensional forms (b=1/4) as well as a fit to a conventional Arrhenius (b=1) behavior (b) Low-field magnetic susceptibility (M/H) collected under $\mu_0 H = 0.1$ T. Dashed line is a fit to a low-temperature Curie-law with the additional of a substantial Pauli-like χ_0 term. Inset shows the magnetic susceptibility measured across the partial freezing transition near 1.5K. For molar magnetic susceptibility, 1 emu/(mol Oe) = $4\pi\ 10^{-6}$ m^3/mol (c) Isothermal magnetization data collected at 1.8 K and 300 K. Dashed line shows a fit using a dominant, linear χ_0 term and a S=1/2 Brillouin function. (d) Isobaric heat capacity $C_p(T)$ collected under a variety of magnetic fields. Inset shows the magnetic entropy extracted via removal of the nonmagnetic analog $NaRhO_2$. (e) Low-temperature heat capacity collected below the crossover at 1.5 K. Power-law fits are shown as solid lines (f) $C_p(T)/T$ plotted as a function of T^2. Solid lines are fits to the model described in the text. Inset shows the evolution of the effective Sommerfeld coefficients under the application of a magnetic field. Error bars represent one standard deviation.

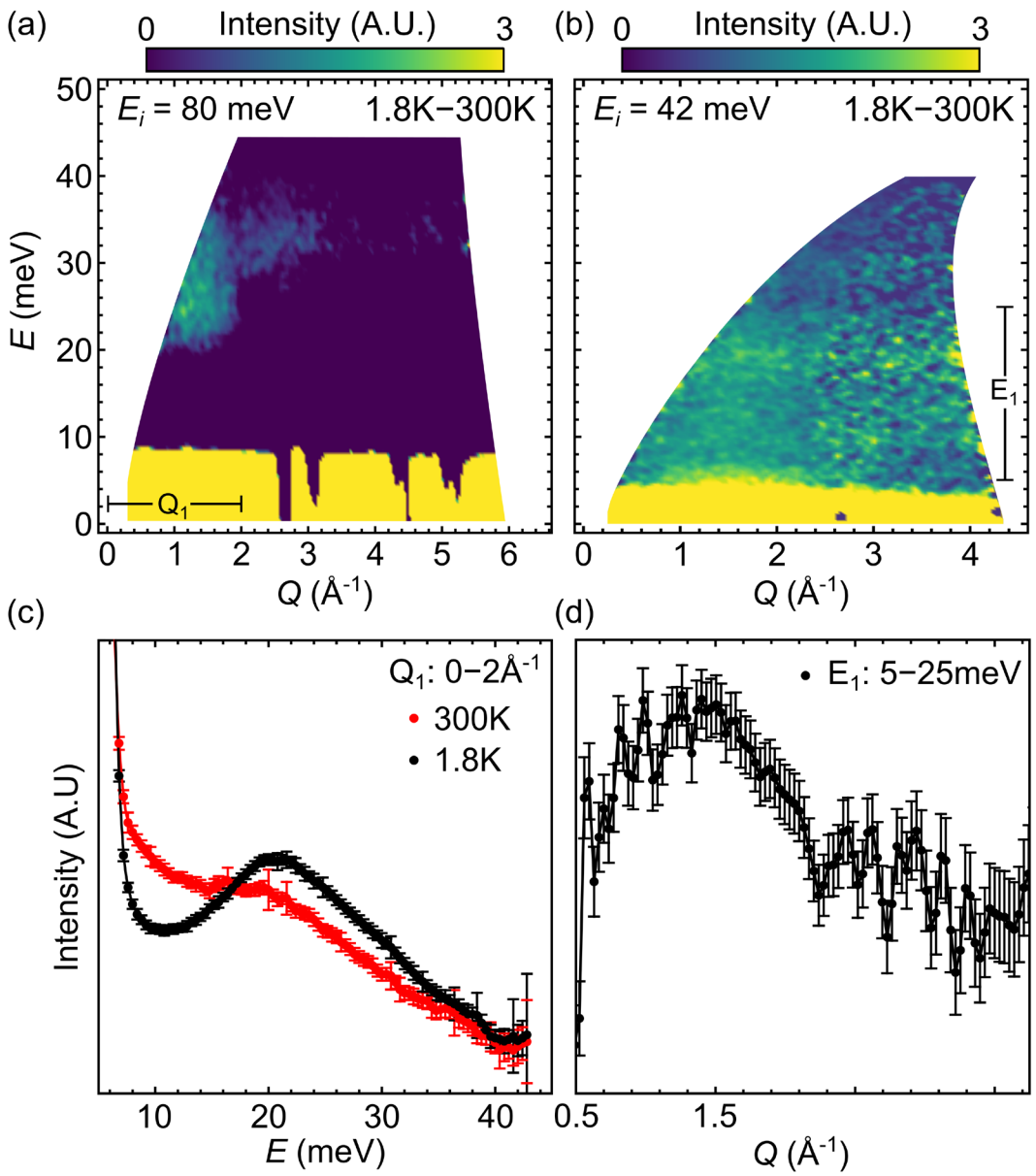

Figure 3: Inelastic neutron scattering data collected on $NaRuO_2$ (a) Energy-momentum map of scattering intensity collected at 1.8 K with the 300 K data directly subtracted. (b) Energy-momentum map of scattering intensity collected at 1.8 K with the 300 K data subtracted after removing the Bose population factor (c) Momentum-averaged energy cut through the temperature subtracted map plotted in panel (a), (d) Energy-averaged momentum cut through the map plotted in panel (b). Error bars represent one standard deviation.

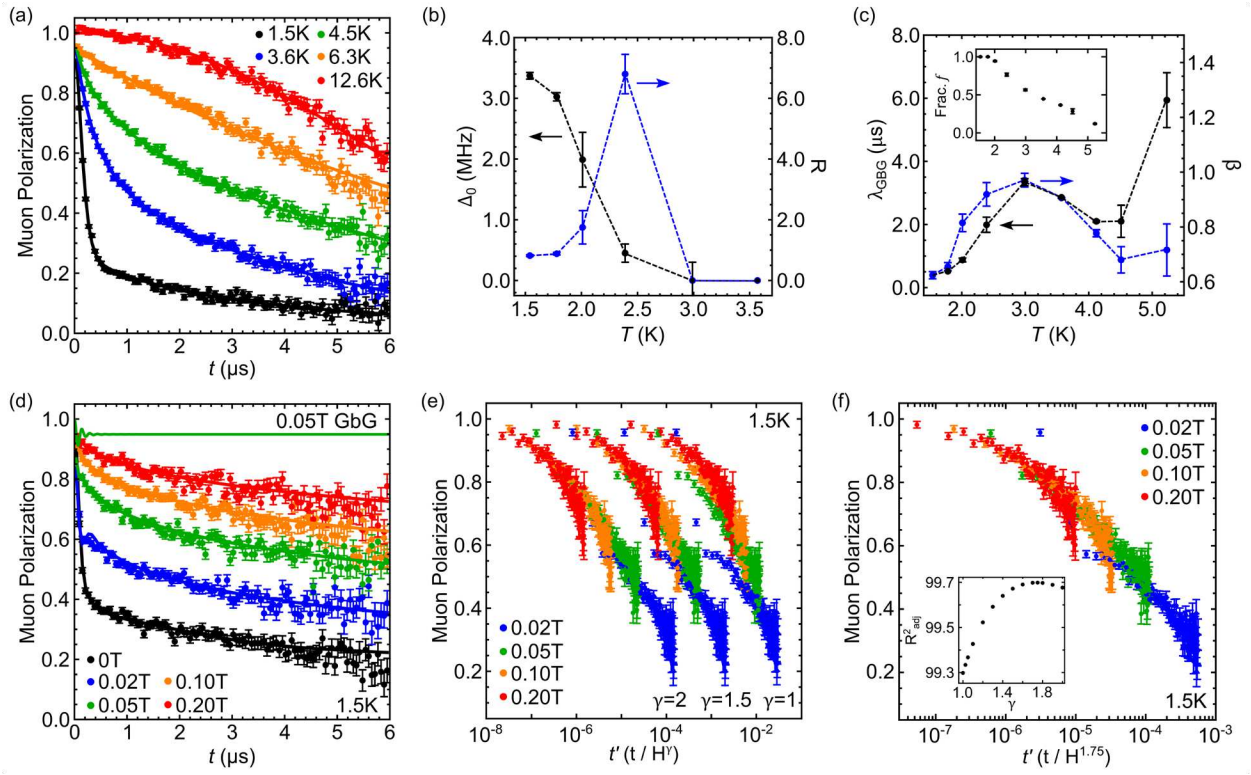

Figure 4: Muon spin relaxation data and analysis (a) Zero-field muon spin relaxation collected for a variety of temperatures. (b), (c) Fit values for the zero-field muon spin relaxation model described in the text. (d) Muon spin relaxation data collected at 1.5 K under a variety of applied longitudinal fields; the solid green curve is the calculated response for a fully static GbG function for the zero-field fit parameters and γ_{GbG} set to 0 at T=1.5 K. (e) 1.5 K muon polarization under varying longitudinal field strengths plotted with different exponents of t/H^{γ}. (f) Scaling collapse of 1.5 K longitudinal field data for an exponent of γ=1.75. Inset: Goodness of fit of the data to a 3rd order polynomial as a function of γ. Error bars represent one standard deviation.

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Figures

Figure 1

Lattice and electronic band structures of $NaRuO_2$ (a) Crystal lattice of $NaRuO_2$ showing the triangular lattice of edge-sharing RuO_6 octahedra arranged in a layered structure separated by planes of Na-ions. The slight trigonal distortion of the RuO_6 octahedra is illustrated. (b) Neutron powder diffraction data collected at 300 K, 1.5 K, 1 K, and 50 mK. Data in the main panel were collected at BT-1 and are overplotted with the refined structural diffraction pattern. The inset shows data collected on HB-1A and reveals no magnetic scattering appearing down to 50 mK. (c) LDA electronic band structure for $NaRuO_2$ using both l and U=1 eV.

Figure 2

Electrical transport, magnetic susceptibility, and heat capacity data characterizing the low temperature properties of $NaRuO_2$ (a) DC resistivity data collected as a function of temperature. Fits to models of variable-range hopping transport are overpotted for both two-dimensional (b=1/3) and three-dimensional forms (b=1/4) as well as a fit to a conventional Arrhenius (b=1) behavior (b) Low-field magnetic susceptibility (M/H) collected under m_0H = 0.1 T. Dashed line is a fit to a low-temperature Curie-law with the additional of a substantial Pauli-like c_0 term. Inset shows the magnetic susceptibility measured across the partial freezing transition near 1.5K. For molar magnetic susceptibility, 1 emu/(mol Oe) = $4\pi\ 10^{-6}$ m^3/mol (c) Isothermal magnetization data collected at 1.8 K and 300 K. Dashed line shows a fit using a dominant, linear c_0 term and a S=1/2 Brillouin function. (d) Isobaric heat capacity $C_p(T)$ collected under a variety of magnetic fields. Inset shows the magnetic entropy extracted via removal of the nonmagnetic analog $NaRhO_2$. (e) Low-temperature heat capacity collected below the crossover at 1.5 K. Power-law fits are shown as solid lines (f) $C_p(T)/T$ plotted as a function of T^2. Solid lines are fits to the model described in the text. Inset shows the evolution of the effective Sommerfeld coefficients under the application of a magnetic field. Error bars represent one standard deviation.

Figure 3

Inelastic neutron scattering data collected on $NaRuO_2$ (a) Energy-momentum map of scattering intensity collected at 1.8 K with the 300 K data directly subtracted. (b) Energy-momentum map of scattering intensity collected at 1.8 K with the 300 K data subtracted after removing the Bose population factor (c) Momentum-averaged energy cut through the temperature subtracted map plotted in panel (a), (d) Energy-averaged momentum cut through the map plotted in panel (b). Error bars represent one standard deviation.

Figure 4

Muon spin relaxation data and analysis (a) Zero-field muon spin relaxation collected for a variety of temperatures. (b), (c) Fit values for the zero-field muon spin relaxation model described in the text. (d) Muon spin relaxation data collected at 1.5 K under a variety of applied longitudinal fields; the solid green curve is the calculated response for a fully static GbG function for the zero-field fit parameters and g_{GbG} set to 0 at T=1.5 K. (e) 1.5 K muon polarization under varying longitudinal field strengths plotted with different exponents of t/H^g. (f) Scaling collapse of 1.5 K longitudinal field data for an exponent of g=1.75. Inset: Goodness of fit of the data to a 3rd order polynomial as a function of g. Error bars represent one standard deviation.

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