Magnetic properties of equiatomic CrMnFeCoNi

Timothy A. Elmslie, Jacob Sartt, Sujely Soto-Medina, Yang Yang, Keke Feng, Ryan E. Baumbach, Emma Zappala, Gerald D. Morris, Benjamin A. Frandsen, Mark W. Meisel, Rémi Dingreville, and James J. Hamlin

1Department of Physics, University of Florida, Gainesville, Florida 32611, USA
2Department of Materials Science and Engineering, University of Florida, Gainesville, Florida 32611, USA
3Center for Integrated Nanotechnologies, Sandia National Laboratories, Albuquerque, New Mexico 87185, USA
4Department of Physics, Florida State University and National High Magnetic Field Laboratory, Tallahassee, Florida 32310, USA
5Department of Physics and Astronomy, Brigham Young University, Provo, Utah 84602, USA
6Centre for Molecular and Materials Science, TRIUMF, Vancouver, British Columbia, Canada V6T 2A3
7Department of Physics and the National High Magnetic Field Laboratory, University of Florida, Gainesville, Florida 32611, USA

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Magnetic, specific heat, and structural properties of the equiatomic Cantor alloy system are reported for temperatures between 5 and 300 K, and up to fields of 70 kOe. Magnetization measurements performed on as-cast, annealed, and cold-worked samples reveal a strong processing history dependence and that high-temperature annealing after cold working does not restore the alloy to a “pristine” state. Measurements on known precipitates show that the two transitions, detected at 43 and 85 K, are intrinsic to the Cantor alloy and not the result of an impurity phase. Experimental and ab initio density functional theory computational results suggest that these transitions are a weak ferrimagnetic transition and a spin-glass-like transition, respectively, and magnetic and specific heat measurements provide evidence of significant Stoner enhancement and electron-electron interactions within the material.

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I. INTRODUCTION

Investigation of high-entropy alloys began in the early 2000s with the work of Cantor, Kim, and Warren [1], who investigated multicomponent alloys of the formula $A_{x}B_{1-x}Al_{10}$, in which $A$ and $B$ each represented a combination of two to four different elements in equiatomic ratios. In this way, they were able to produce stable combinations of up to seven different elements. The phrase “high-entropy alloy,” however, was not introduced until 2004, in the work of Yeh et al. [2], who used it to describe compounds consisting of five or more elements. These alloys are of interest partly due to the extremely large number of new alloy systems which fall into the category of high- or medium-entropy alloys [2]. Furthermore, prior investigations have revealed a number of alloys with intriguing properties such as high hardness and resistance to anneal softening [2], shape memory effects [3,4], and superconductivity [5]. Cantor alloys also have potential applications due to soft magnetism [6] and other tunable magnetic properties [7].

The specific combination of chromium (Cr), manganese (Mn), iron (Fe), cobalt (Co), and nickel (Ni) that would come to characterize the “Cantor alloy” was first identified in the 2004 work of Cantor, Chang, Knight, and Vincent [8], who produced alloys of 16 and 20 different elements in equal proportions through induction melting. The resulting compound was unsurprisingly multiphase, but the predominant phase was especially rich in Cr, Mn, Fe, Ni, and Co. In the same work, this CrMnFeCoNi compound was identified as face-centered cubic (fcc), and the ability of this material to dissolve large amounts of elements such as niobium (Nb), titanium (Ti), and vanadium (V) was noted.

Much work has been done regarding the mechanical properties of Cantor alloys [8–13], as well as the magnetic properties of similar compounds [6,7,14–23], but fewer studies have examined the magnetic properties of the original CrMnFeCoNi alloy [24–28]. Of its component elements, Cr and Mn are antiferromagnetic with $T_N = 311 K$ [29] and $T_N = 100 K$ [30] respectively, while Fe, Co, and Ni are ferromagnetic with $T_C = 1043 K$, $T_C = 1394 K$, and $T_C = 631 K$, respectively [31]. However, the magnetic transitions in the equiatomic Cantor alloy appear at much lower temperature. Jin et al. [24] identified a peak in the magnetization of CrMnFeCoNi at 25 K and suggested that it could be either an antiferromagnetic transition or a spin-glass transition. Schneeweiss et al. [26], however, found two transitions in the magnetization: a spin-glass transition at 93 K, and a ferromagnetic transition at 38 K. They were able to identify the 38 K transition as ferromagnetic by magnetic hysteresis loops after cooling in field, while the nature of the spin-glass transition was confirmed by magnetic moment relaxation measurements. Despite identifying these magnetic characteristics, the data of Schneeweiss et al. do not afford additional quantitative analysis since the magnetization results are only reported in units of emu, rather than in a generic manner like emu g$^{-1}$. Consequently, the identification of the atoms
and/or atomic morphology generating the magnetic response is not quantitatively addressed, and this work provides insights about this important issue.

Kamarád et al. [27] investigated the magnetic properties of the equiatomic Cantor alloy between ambient pressure and 1 GPa, observing that increasing pressure decreased magnetization slightly. Using Curie-Weiss fitting, they found evidence of strong antiferromagnetic interactions, which were conjectured to be responsible for the observed small magnetization values as well as the linear field dependence of the magnetization. In contrast, while Schneeweiss et al. found ferromagnetic-type behavior in magnetic hysteresis loops after cooling in an applied magnetic field, Kamarád et al. were unable to replicate these results, seeing a change of only 0.4 emu g\(^{-1}\) in magnetization after field cooling. These observations led Kamarád et al. to disagree with Schneeweiss et al. on the nature of the magnetic transitions within the compound, identifying ferrimagnetic order below 85 K and magnetic cluster-glass behavior below 43.5 K.

This work reports a quantitative analysis of the magnetic properties of the equiatomic Cantor alloy CrMnFeCoNi based on a combination of compositional and structural characterization, magnetization studies, Hall effect measurements, muon spin relaxation (\(\mu\)SR) results, specific heat studies, and \textit{ab initio} density functional theory (DFT) calculations. The magnetic response of as-cast, annealed, and cold-worked samples revealed a strong history dependence, which is described and contrasted with the magnetic properties of precipitates known to form in Cantor alloys. All of our results lead to the identification of the magnetic signatures as intrinsic characteristics of the equiatomic Cantor alloy CrMnFeCoNi, with weak ferrimagnetic order near 43 K, a spin-glass-like fingerprint near 85 K, and a sizable temperature-independent response. This last contribution was established by Curie-Weiss analysis of the magnetic susceptibility data and further probed by Hall measurements to determine the expected size of the Pauli paramagnetic contribution to the susceptibility. Combined with an interpretation of the specific heat data and DFT results, our work provides evidence of a sizable Stoner enhancement and electron-electron interactions.

II. METHODS

A. Materials synthesis and history

Samples were synthesized by combining stoichiometric amounts of elemental Cr, Mn, Fe, Co, and Ni and melting them together in an Edmund Bühler MAM-1 compact arc melter to produce as-cast samples. The Cr, Mn, Fe, and Ni elements were sourced from Alfa Aesar, while Co was purchased from Cerac. The Cr used for synthesis was 99.995% pure, while the Co was 99.5% pure. All other elements were 99.95% pure. Each sample was melted five times, flipping over between each melt to improve sample homogeneity. Samples measured immediately after arc melting are referred to as “as-cast.” Annealed samples were made by sealing as-cast samples in quartz tubes under Ar atmosphere and homogenizing them at 1100 °C for 6 days, after which the tube containing the samples was quenched in water. Samples measured following this step are called “anneal A.” The piece cut from the as-cast boule and annealed in ambient atmosphere at 700 °C for 1 h is referred to as “oxidized.” Some samples from the anneal A batch were cold worked by flattening them in a hydraulic press a total of three times using a pressure of approximately 0.5 GPa, folding them in half between each flattening step. These samples are known as “cold-worked.” After cold working, the samples were reannealed in quartz tubes under argon (Ar) atmosphere. A portion were annealed at 700 °C for 1 h, while others were again subjected to 1100 °C for 6 days, referred to as “anneal B” and “anneal C,” respectively.

B. Compositional and structural analysis

The microstructure of the sample was characterized using a Tescan MIRA3 scanning electron microscope (SEM) with an energy-dispersive x-ray (EDX) detector at 20 kV. Prior to SEM measurements, samples were mounted in resin epoxy while first polished using 600-grit, then 800-grit SiC paper. This surface was further polished using alcohol-based lubricant and diamond paste of varying particulate sizes in steps of 6, 3, and 1 \(\mu\)m, followed by 0.05-\(\mu\)m master polishing using water-free colloidal silica suspension. The crystal structure of the sample was investigated using a Panalytical Xpert powder x-ray diffractometer (XRD) with a copper-radiation source at an accelerating voltage of 40 kV and the electron current of 40 mA, over a 2\(^\circ\) range from 40\(^\circ\) to 120\(^\circ\). Powder samples were prepared using a mortar and pestle. Quantitative composition data were obtained by electron probe microanalysis (EPMA) using a CAMECA SXFiveFE instrument operating with an accelerating voltage of 15 kV and a probe current of 20 nA.

C. Magnetic and magnetotransport studies

A Quantum Design Magnetic Property Measurement System (MPMS) was used to take magnetization data. Small pieces ranging from a few milligrams to a few hundred milligrams were cut from larger samples using an Allied 3000 low-speed saw to minimize unintentional working of the samples. Each sample was secured in a gel capsule inside a plastic straw for measurement. In order to obtain magnetization versus temperature data, samples were cooled under zero field to a temperature of 5 K, after which field was applied and data were recorded while warming. This process produced the datasets labeled zero-field cooled (ZFC). The applied field was held constant while the temperature warmed to 300 K and subsequently cooled to 5 K again. Then data were recorded while warming to produce field-cooled warming (FCW) datasets.

Resistivity and Hall measurements were performed in a Quantum Design Physical Property Measurement System (PPMS) at 300 K. For these measurements, an Allied 3000 low-speed saw was used to cut an 8 \(\times\) 8.4 mm\(^2\) flat square sample from the arc-melted boule. Then four notches, one on each side, were cut into it with the same device to produce a rough cloverleaf shape for the purposes of Van der Pauw measurements. To increase the signal-to-noise ratio, the sample was first sanded on a custom jig to a thickness of 0.2 mm, then polished with 800-grit silicon carbide paper to a final
thickness of 0.16 mm. This sample was placed on a PPMS transport puck with a piece of cigarette paper between sample and puck for insulation, while N grease provided thermal contact. Each “leaf” of the cloverlike shape of the sample was connected to one of the puck solder pads using platinum wire affixed to the sample with silver paint.

D. Muon spin relaxation experiments

Muon spin spectroscopy measurements, or μSR, were conducted to probe the spin dynamics and provide sensitivity to the volume fraction of the various magnetic phases in equiatomic Cantor alloy. These experiments were conducted at TRIUMF Laboratory in Vancouver, Canada, using the LAMPF spectrometer on the M20D beam line. Positive muons with 100% initial spin polarization were implanted one at a time in the sample. After implantation, the muon spin underwent Larmor precession in the local magnetic field at the muon site, which consists of the vector sum of the internal magnetic field due to electronic and/or nuclear dipolar moments and any externally applied magnetic field. After a mean lifetime of 2.2 μs, the muon decays into two neutrinos and a positron, with the latter being emitted preferentially in the direction of the muon spin at the moment of decay. Two detectors placed on opposite sides of the sample record positron events as a function of time after muon implantation, yielding the μSR asymmetry \( a(t) = [N_1(t) - N_2(t)]/[N_1(t) + N_2(t)] \), where \( N_1(t) \) and \( N_2(t) \) are the number of positron events recorded at time \( t \). This quantity is proportional to the projection of the muon ensemble spin polarization along the axis defined by the positions of the detectors, from which information about the local magnetic field distribution in the sample can be inferred [32]. The sample was mounted on a low background copper sample holder, and the temperature controlled using a helium gas flow cryostat. The μSR data were analyzed using the open-source programs MUSRFIT [33] and BEAMS [34], which yield consistent results. The correction parameter \( \alpha \) was determined according to standard practice from a weak transverse field measurement in the paramagnetic state [35].

E. Specific heat measurements

Alloy specific heat was measured in a PPMS with the attached PPMS heat capacity option in order to obtain the Debye temperature and the approximate magnetic entropy of the sample. The sample used for specific heat measurements was annealed at 1100 °C for 6 days before being cut to a mass of a few milligrams using an Allied 3000 low-speed saw to minimize sample deformation and strain. After cutting, this piece was placed on the sample platform of a Quantum Design heat capacity puck using a small amount of Apeizon N grease to adhere the sample and provide thermal contact before being inserted into the PPMS for measurement.

F. DFT calculations

Spin collinear DFT simulations were performed to investigate exchange splitting between spin states and its effect on ferromagnetism within the system. For modeling, the Vienna Ab initio Simulation Package (VASP) [36–38] was used, in which the electronic wavefunctions were modeled through plane waves and projector-augmented wave (PAW) pseudopotentials [39,40]. Exchange-correlation effects were treated within the generalized gradient approximation (GGA) according to the parametrization of Perdew, Burke, and Ernzerhof (PBE) [41], while the individual pseudopotentials used to model each species were chosen so that only the outermost \( s \) and \( d \) valence electrons were explicitly included.

The alloy atomic structure was modeled using supercells containing 108 atoms, representing a \( 3 \times 3 \times 3 \) transformation of the conventional four-atom fcc unit cell. Special quasirandom structures (SQSs) [42,43], built using the openly available Alloy-Theoretic Automated Toolkit (ATAT) [44], were used to replicate random atomic ordering among lattice sites. To ensure the most accurate description of properties of the randomly ordered solid solution structure, four separate 108-atom SQS supercells were modeled at the equiatomic composition, each using a \( 5 \times 5 \times 5 \) gamma-entered \( k \)-point mesh. For all simulations, electronic convergence was met when the total energy of the system fell below \( 1.0 \times 10^{-6} \) eV, while ionic convergence was met when forces on all atoms fell below \( 20 \times 10^{-3} \) eVÅ\(^{-1} \). Lastly, a plane-wave energy cutoff of 400 eV was found to sufficiently minimize the total energy, lattice constants, magnetic moments, and fluctuations in the density of states (DOS) when used with a Gaussian smearing method and a smearing width of 0.01 eV.

Lastly, the experimental determination of the magnetic entropy requires one to know the magnetic contribution to the specific heat capacity. This is commonly achieved by using as a reference value the experimental specific heat of a similar but nonmagnetic alloy that exhibits the same crystal structure. In this work, we were instead able to employ density functional perturbation theory (DFPT) to calculate the specific heat for a nonmagnetic version of the equiatomic Cantor alloy (i.e., modeled without the effects of spin polarization). This calculation was performed using the open-source package PHONOPY [45], in which the specific heat at constant volume \( C_V^{\text{DFT}} \) is calculated through the harmonic approximation as a function of phonon dispersion frequencies \( \omega \), according to

\[
C_V^{\text{DFT}} = \sum_{\mathbf{q}, v} k_B \left( \frac{\hbar \omega_{\mathbf{q}, v}}{k_B T} \right)^2 \exp \left( \frac{\hbar \omega_{\mathbf{q}, v}}{k_B T} \right) \left[ \exp \left( \frac{\hbar \omega_{\mathbf{q}, v}}{k_B T} \right) - 1 \right]^{-2},
\]

where \( T \) is the temperature, \( \omega_{\mathbf{q}, v} \) is the phonon frequency at wave vector \( \mathbf{q} \) and band index \( v \), and \( k_B \) and \( \hbar \) respectively, are the Boltzmann and reduced Planck constants.

III. EXPERIMENTAL RESULTS

A. Microstructural analysis

A representative example of the microstructure of the Cantor alloy samples is shown in Fig. 1 for the anneal A sample. The microsized pores observed in the SEM image are considered to be casting porosity. Additionally, EDX analysis identified a small number of particles as oxide rich in Cr and Mn. The sample consists of a single fcc phase according to the XRD pattern, and no peak was identified as corresponding to the oxide particles. The existence of such (Cr, Mn)-rich particles in as-cast and homogenized Cantor alloys has been reported in previous studies [46–48]. Furthermore, the XRD
FIG. 1. SEM image of homogenized Cantor alloy (anneal A). A small number of particles (black spots identified by arrows) were determined by EDX to be oxide rich in Cr and Mn. The XRD data indicated the sample is single phase and all peaks could be indexed as fcc.

data did not contain any peaks associated with inclusions [49], which, due to their small volume fraction, are not considered to significantly influence the properties reported later in this work. Sample composition was confirmed via an EPMA measurement of the as-cast sample, and the results are presented in Table I.

B. Magnetization of homogenized CrMnFeCoNi

A magnetization-versus-temperature plot for a sample of equiatomic Cantor alloy at different temperatures from 5 to 300 K is displayed in Fig. 2. All of the displayed curves remain roughly linear across the measured range of −70 to 70 kOe with magnitudes of only a few hundredths of a bohr magneton, demonstrating that the compound is far from magnetic saturation. The only feature that clearly distinguishes one curve from another is the temperature dependence as indicated by different slopes for different temperatures. The 300 K dataset has the lowest slope, and slope increases as temperature decreases until 40 and 45 K, which largely overlap one another. As temperature falls below 40 K, the slope begins to decrease again.

FCW magnetization versus temperature data from a sample of anneal A are displayed in Fig. 3 at fields from 0.1 to 20 kOe, while the inset compares both ZFC and FCW data at 0.1 kOe. Magnetization plots are presented in units of emu mol\(^{-1}\) Oe\(^{-1}\) to simplify estimation of effective moment, \(p_{\text{eff}}\), which can be written as

\[
p_{\text{eff}} = 2.82 C_m^{1/2},
\]

where \(p_{\text{eff}}\) is in units of Bohr magnetons and \(C_m\) is the Curie constant in units of emu K mol\(^{-1}\) Oe\(^{-1}\) [50]. In this notation,

TABLE I. Composition of the as-cast equiatomic Cantor alloy, as determined by EPMA measurement.

| Element | at. % |
|---------|-------|
| Cr      | 20 ± 1 |
| Mn      | 19 ± 2 |
| Fe      | 20 ± 1 |
| Co      | 20 ± 1 |
| Ni      | 20 ± 1 |

FIG. 2. Magnetization versus field data on equiatomic Cantor alloy at a variety of temperatures between 5 and 300 K. Inset shows the same data at low field, between −6 and 6 kOe. The magnetization remains small to the highest measured fields.

FIG. 3. Magnetic susceptibility versus temperature data on equiatomic Cantor alloy under various applied magnetic fields. The main plot shows only field-cooled warming (FCW) data, while the inset compares zero-field cooled (ZFC) and FCW data at 0.1 kOe.
mol\textsubscript{atom} denotes moles of atoms, as opposed to, for example, moles of formula units or moles of magnetic ions, and emu is defined as erg G\textsuperscript{-1}, as reported by the Quantum Design MPMS. At 0.1 kOe, two transitions are clearly seen: a steplike anomaly at 43 K, and a peak at 85 K. As field increases, the 43 K transition is rapidly suppressed, becoming notably smaller at 1 kOe and vanishing completely at 5 kOe and above. The higher temperature transition is less affected by the increase in field. At an applied field of 20 kOe, the higher temperature transition is slightly smaller, slightly broadened, and shifted down in temperature from 85 to 75 K compared to the 0.1 kOe curve. The small size of the 43 K transition is notable, which led us to examine the possibility that it could derive from one of the known impurity phases [47,51,52].

C. Effects of plastic deformation

Since the Cantor alloy is known to form precipitates as a result of deformations and moderate temperature anneals [47,51,52], investigation of the processing history dependence of the compound is necessary in order to understand the origin of the magnetic behavior. The 43 K transition in particular is notably small, with step size of approximately $1 \times 10^{-5} \mu_B$ atom\textsuperscript{-1} in fields of 0.1 and 1 kOe, suggesting that it may originate from a magnetic secondary phase which comprises a proportionally small amount of the sample. This hypothesis is tested through magnetization measurements performed on samples which have undergone different cold-working and annealing processes. The data are plotted in Fig. 4, in which each curve represents a different processing history, as described in the Methods section. In Fig. 4(a), the as-cast sample shows a large 43 K transition relative to the other samples, but notably lacks a peak at higher temperature. Annealing the as-cast material at 1100 °C produces the anneal A sample, in which the 45 K signature is significantly smaller and sharper than the corresponding feature in the as-cast sample. Additionally, a sharp peak appears at 85 K in the anneal A data. Further changes of the magnetic response are observed after cold working the anneal A sample in a hydraulic press (cold-worked), and subsequently, annealing at 700 °C for 1 h (anneal B), and at 1100 °C for 6 days (anneal C). These datasets demonstrate that cold working reduces the size of the 43 K transition and pushes the higher temperature transition to lower temperatures while increasing its size. However, annealing after cold working, even for a brief duration, results in a 43 K transition larger than what appears in the cold-worked samples but smaller than that of the arc-melted as-cast samples. In addition, in moving from the as-cast to the annealed samples, the higher temperature transition is revealed, even if it remains smaller than in the cold-worked sample. This strong processing dependence also explains differences between the results of our measurements and those of others [24,26,27].

D. Impurity contribution to magnetic properties

Having investigated the effects of different processing methods on the equiatomic Cantor alloy itself, the next step was to synthesize and measure the magnetism of the precipitates, shown in Fig. 5, with the annealed equiatomic, NiMn, and NiMn\textsuperscript{+}(Cr,Fe,Co)\textsubscript{ε} on the left y axis of Fig. 5(a) and FeCo and FeCo\textsuperscript{+}(Cr,Mn,Ni) on the right axis. Despite “pristine” annealed sample. The inset plot shows a detailed view of low-field data below 6 kOe. Anneal A reveals the lowest level of hysteresis, while the cold-worked and anneal C samples show the highest. Larger hysteresis may be indicative of the emergence of ferromagnetic secondary phases.
FIG. 5. (a) Comparison of magnetic susceptibility versus temperature data from FeCo and NiMn precipitate compounds as well as an oxidized equiatomic compound and anneal A. FeCo-based compounds exhibit comparatively large magnetization and are measured on the right axis for clarity. (b) Comparison of magnetic susceptibility data from the Cr-rich precipitate compound and anneal A. The “+(Cr,Fe,Co)”, “+(Cr,Mn,Ni)”, and “+(Mn,Fe,Co,Ni)” suffixes indicate the presence of small amounts of other Cantor alloy elements consistent with the precipitate compositions observed in the works of Otto et al. [47], Schuh et al. [51], and Li et al. [52]. Closed circles represent ZFC data, while open circles represent FCW data. All data are measured in emu per moles of atoms per Oe.

synthesizing and annealing in Ar atmosphere, oxidation of the alloy is also considered, and data from the oxidized sample are also included on the left axis of Fig. 5(a). Measurements performed on the Cr-rich precipitate reveal magnetization values significantly higher than those of the NiMn compounds and significantly lower than the FeCo compounds, such that the Cr compound data do not fit easily alongside either. For clarity, this curve has been separated into Fig. 5(b), where it is compared to anneal A independently. The suffixes of “+(Cr,Fe,Co)”, “+(Cr,Mn,Ni)”, and “+(Mn,Fe,Co,Ni)” indicate the presence of small amounts of other Cantor alloy elements added to match the compositions of precipitates reported by Otto et al. [47], Schuh et al. [51], and Li et al. [52]. The investigated precipitates lack any signs of a transition near 43 K, suggesting that this transition is, in fact, an intrinsic aspect of the Cantor alloy system. While the 85 K transition is present in the oxidized curve, its size is not significantly changed relative to the anneal A curve. Furthermore, this transition is also not present in any of the precipitate curves, suggesting that it is also intrinsic to the Cantor alloy.

E. Muon spin relaxation

Key experimental results from the μSR measurements of a sample from anneal A are shown in Fig. 6. Figure 6(a) displays asymmetry spectra collected at representative temperatures between 2 and 153 K. At the highest temperature studied, a gentle relaxation with a Gaussian-like nature is observed. Applying a field of 1 kOe parallel to the initial muon
spin polarization recovers approximately 75% of the asymmetry (data not shown), indicating that the relaxation at this temperature is due primarily to weak, random dipolar fields from nuclear moments [53]. As the temperature is lowered, the relaxation increases and becomes more exponential in nature, characteristic of magnetic fluctuations from electronic spins. Between 87 and 82 K, the initial asymmetry drops rapidly from ~0.24 to ~0.135 and then further to 0.08 (i.e., 1/3 the total initial asymmetry) at 77 K. This “1/3 tail” is the hallmark of a polycrystalline material with static magnetism throughout the full sample volume. No coherent oscillations of the asymmetry are observed in the “2/3 component,” indicating very large fields and/or a broad distribution of field strengths at the muon stopping sites [53]. As the temperature is lowered further, the relaxation of the 1/3 component gradually slows until almost no relaxation remains at 2 K, consistent with magnetic fluctuations freezing out. Qualitative inspection of the spectra therefore confirms the presence of a sharp and uniform transition throughout the full sample volume with an onset temperature between 82 and 87 K. No pronounced change in the spectra is observed around 43 K, which demonstrates the transition at this temperature cannot be due simply to a minority phase in the sample; otherwise a further drop of the asymmetry would be observed below 43 K.

To gain more quantitative insight, least-squares fits to the asymmetry data were performed using the stretched exponential function \( a(t) = a_0 e^{-(t/\lambda)^\beta} \), where \( a(T) \) is the temperature-dependent asymmetry, \( a_0 \) is the initial asymmetry at \( t = 0 \), \( \lambda \) is the relaxation rate, and \( \beta \) is the exponential power. This type of stretched exponential function is phenomenological, employed to model a continuous distribution of relaxation rates [54]. The best-fit asymmetry curves agree well with the data, as seen by the solid curves in Fig. 6(a). The volume fraction of the sample that exhibits static magnetism can be determined from the fitted asymmetry values as \( f(T) = (a_{0\text{max}} - a_0(T))/(a_{0\text{max}} - a_{0\text{LT}}) \), where \( a_0(T) \) is the best-fit value of \( a_0 \) at temperature \( T \), and \( a_{0\text{max}} \) is the maximum value of \( a_0 \) across all measured temperatures, and \( a_{0\text{LT}} \) is the average value of \( a_0 \) for the low-temperature data (\( T < 70 \) K). As seen in Fig. 6(b), \( f(T) \) transitions rapidly from zero to 1 in a small temperature window centered around 82 K, indicating that the sample undergoes a highly uniform magnetic transition. Evidence of this transition is also shown in Fig. 6(c), which displays the relaxation rate \( \lambda \) and the exponential power \( \beta \) as a function of temperature. The prominent peak in \( \lambda \) centered around 82 K is evidence of critical spin dynamics as the temperature decreases toward the transition, as is observed in canonical spin glasses and continuous phase transitions [35,55]. The exponential power \( \beta \) is \( \sim 1.5 \) at high temperature where relaxation from nuclear dipolar fields dominates, but it decreases steadily as the temperature is lowered and electronic spin fluctuations become more prominent. At the transition temperature and below, \( \beta \) converges to values between 0.45 and 0.55 when left fully unconstrained, and for consistency, \( \beta \) was set to 0.5 for the spectra collected at 82 K and below. This value of \( \beta \) is expected when the system exhibits multiple relaxation channels and/or spin fluctuation rates [56], which is unsurprising in this highly disordered alloy. Similar values were also observed in the entropy-stabilized antiferromagnetic oxide (Mg,Co,Ni,Cu,Zn)O [57].

\[ C = \gamma T + \beta \rho \delta T^3, \]  

(3)

the expression for the low-temperature specific heat of a metal [60]. In this equation, the \( \gamma T \) term represents the electronic component of the specific heat, while \( \beta \delta T^3 \) is the lattice specific heat and is related to the Debye temperature. Fitting to Eq. (3) gives

\[ \gamma = 15.3(2) \text{ mJ mol}^{-1} \text{ K}^{-2}, \]

\[ \beta = 0.025(3) \text{ mJ mol}^{-1} \text{ atom}^{-1} \text{ K}^{-4}. \]

The magnetic contribution to the specific heat for a ferromagnet at low temperature can be represented by a term \( \delta T^3/2 \), while for an antiferromagnet the contribution goes like \( T^3 \) [61–66]. Including a \( T^{3/2} \) term in the fitting equation results in a negligibly small \( \delta \) coefficient and does not significantly affect the other fitting values, suggesting that ferromagnetic contributions to the low-temperature specific heat are not significant. In the antiferromagnetic case, an estimate of the magnetic contribution to the low-temperature specific heat is
complicated by the fact that the expected temperature dependence ($T^3$) is the same as that of the lattice, so that the two contributions cannot be straightforwardly disentangled.

G. Hall effect and carrier density

Measurements of Hall voltage as a function of magnetic field permit the estimation of carrier density according to

$$V_H = \frac{IB}{ned},$$

(4)

where $I$ is the current, $B$ is the magnetic field perpendicular to the current, $e$ is the electron charge, and $d$ is the sample thickness. The carrier density can, in turn, provide insight into the magnetism of the alloy as carried out in Sec. IV C. Results of Hall measurements are shown in Fig. 8, including a linear fit with a slope of $1.4(4) \times 10^{-4} \text{nV Oe}^{-1}$. Using this slope in combination with Eq. (4), a carrier density $n = 2.9(8) \times 10^{22} \text{ cm}^{-3}$ can be calculated. This result is consistent with the carrier density of Al$_x$CrFeCoNi reported by Kao et al. [16], as well as with that of a typical metal ($\sim 1 \times 10^{22}$ to $1 \times 10^{23} \text{ cm}^{-3}$) [67,68].

IV. DISCUSSION

Experimental results reported in this work demonstrate the effects of processing on the two observed transitions in the equiatomic Cantor alloy as well as the intrinsic nature of these transitions. However, the magnetic properties can be examined in more detail, and information on the nature of magnetism in this sample can be extracted from the presented experimental data. This section begins with Curie-Weiss analysis of the magnetic susceptibility which reveals a significant constant offset. Analysis which combines specific heat data, computational results, and Hall measurements points to Stoner enhancement as the origin of the offset. The Stoner enhancement parameter $Z$ is calculated through three different methods which obtain a $Z$ value between 0.92 and 0.97, suggesting that this compound is on the cusp of magnetic order. This result is perhaps unsurprising given the observed magnetic transitions and the strong magnetic ordering seen in the Cantor alloy’s constituent elements, but more notable is the small size of the magnetic transitions, which may be due to the presence of antiferromagnetic interactions, as evidenced by DFT results. These findings are consistent with previous work on a similar compound, Fe$_{40}$Mn$_{40}$Cr$_{10}$Co$_{10}$ [69].

A. Modified Curie-Weiss fitting

When using a Curie-Weiss expression to estimate the effective moment of the anneal A sample at temperatures above the 85 K transition, reasonable fits required a modified expression written as

$$\chi(T) = \chi_o + \frac{C_m}{T - \theta},$$

(5)

where $C_m$ is the Curie-Weiss constant, $\theta$ is the Curie-Weiss temperature, and $\chi_o$ is a constant that represents the summation of multiple temperature-independent terms such as Pauli paramagnetism, Van Vleck paramagnetism, Landau diamagnetism, and core diamagnetism [70–74]. Fits which excluded the $\chi_o$ term did not accurately fit the data, and the origin of this term is analyzed in greater detail in Secs. IV C and IV D. The resulting fit is compared with experimental data in Fig. 9, where it is plotted as susceptibility times temperature for greater readability across the entire temperature range, and the resultant value for each parameter is listed in Table II. This fitting equation and its results differ notably from those of Kamarád et al. [27], who used a nonmodified Curie-Weiss equation to obtain a Curie-Weiss temperature of $\theta = -210 \text{ K}$ and an effective moment of $p_{\text{eff}} = 2.71 \mu_B/\text{f.u.}$, using their formula of $\text{Cr}_{0.205}\text{Mn}_{0.20}\text{Fe}_{0.205}\text{Co}_{0.199}\text{Ni}_{0.191}$. Given the strong history dependence of the Cantor alloy, this
TABLE II. Constant offset $\chi_0$, Curie-Weiss constant $C_m$, Curie-Weiss temperature $\theta$, and effective moment $p_{\text{eff}}$ from modified Curie-Weiss fit, Eq. (5). The value of $p_{\text{eff}}$ was calculated using Eq. (2) and the value of $C_m$.

| Variable | Value | Units |
|----------|-------|-------|
| $\chi_0$ | $5.4 \times 10^{-4}$ | emu mol$^{-1}$Oe$^{-1}$ |
| $C_m$    | 0.13  | emu K mol$^{-1}$Oe$^{-1}$ |
| $\theta$ | $-16.1$ | K |
| $p_{\text{eff}}$ | 1.01 | $\mu_B$ atom$^{-1}$ |

The discrepancy may be a result of the differing processing methods, since Kamarád et al. cold-rolled their samples after casting and then annealed at 1173 K for 1 h. Our investigation of Cantor alloy processing history dependence shows that cold working will increase magnetization, and subsequent high-temperature annealing will not restore the sample to a “pristine” state.

B. Specific heat and entropy

The fit in Fig. 7 provides the parameter $\beta$, which is the coefficient of the phonon contribution to the specific heat. The value of $\beta$ can be used to calculate the Debye temperature $\Theta_D$ according to

$$\beta = \frac{12\pi^4 N_A k_B}{5 \Theta_D^3},$$

in which $N_A$ is Avogadro’s number [75]. This calculation gives $\Theta_D = 427$ K, not far from the Debye temperatures of some of the Cantor alloy’s component elements [76]. These values are compared in Table III.

A computational estimate of the lattice component of the specific heat was produced through DFT calculation in the absence of spin-polarization effects. This estimate, denoted as “DFT $C_{\text{lattice}}$,” is shown in the inset of Fig. 10, where it is compared with experimentally obtained specific heat data. Plotting the DFT-calculated lattice specific heat as $C/T$ versus $T$ (not shown) produces a peak at approximately 105.3 K. For the Debye function divided by temperature, the maximum occurs at $0.28 \Theta_D$. Applying this criterion to the DFT specific heat indicates $\Theta_D = 376$ K, which is consistent with the Debye temperature estimates based on the measured data.

TABLE III. Debye temperatures of the equiatomic Cantor alloy and its component elements. Elemental Debye temperatures are obtained from Kittel [76].

| Compound | $\Theta_D$ (K) |
|----------|--------------|
| CrMnFeCoNi, Eq. (6) | 427 |
| CrMnFeCoNi, DFT | 376 |
| Cr | 630 |
| Mn | 410 |
| Fe | 470 |
| Co | 445 |
| Ni | 450 |

FIG. 10. An estimate of the combined electronic and magnetic specific heat produced by subtracting the DFT-calculated lattice contribution shown in Fig. 7. The dashed line represents a linear fit to high-temperature data. The inset compares the measured specific heat to the DFT-calculated result.

Additionally, by subtracting the lattice specific heat from the total measured specific heat, an estimate of the combined magnetic and electronic components is produced. We note that the measurements were performed at constant pressure, while the DFT calculations provide the specific heat at constant volume. However, the difference between $C_p$ and $C_v$ is expected to be of order 1% or less in the relevant temperature regime [60], so that the difference can be ignored in the following analysis.

The results of this subtraction are displayed in the main panel of Fig. 10, alongside a linear fit to the high-temperature data with slope 10.6 mJ mol$^{-1}$atom$^{-2}$ K$^{-2}$. This slope is notably similar to the value of $\gamma$ obtained via a low-temperature fit to the measured specific heat displayed in Fig. 7. Isolating the magnetic component is complicated by potential changes in $\gamma$ across the measured temperature range, similar to the temperature-dependent effective mass observed in zinc [77], such that a linear electronic component cannot simply be subtracted from the data in Fig. 10. Even so, Fig. 10 can provide a rough estimate of the magnetic entropy and therefore the number of spins involved in a magnetic transition. Performing this calculation for our specific heat data provides a value of 0.096 $R \ln 2$ at approximately 50 K and 0.23 $R \ln 2$ at 97 K. In both cases the value is significantly suppressed below that expected for full local moments, suggesting itinerant magnetism.

C. Effective mass and carrier density

The fit shown in Fig. 7 also provides an estimation of the electronic component of specific heat through the coefficient $\gamma$. This value can be used to calculate the effective carrier
The spin-up and spin-down occupations at the Fermi energy, normalized to a per-atom occupation, is presented in Fig. 11. These DOS curves represent a combination of the individual DOS plots from each DFT simulation, normalized for the total number of atoms.

mass $m^*$ according to

$$m^* = \gamma \gamma_0,$$

in which $m_e$ is the mass of a free electron [60]. The variable $\gamma_0$ is a theoretical value calculated from the electronic DOS and can be expressed as

$$\gamma_0 = \frac{\pi k_B^2 g(E_F)}{3},$$

in which $k_B$ is the Boltzmann constant and $g(E_F)$ is the DOS in the $d$ band at the Fermi level, $E_F$ [78]. In metals such as the Cantor alloy studied here, the $s$ and $p$ bands are broad near the Fermi level and thus contribute very little to the overall electron occupation. In other words, only the $d$ band needs to be taken into account [79]. The spin-separated projected DOS of the $d$ band, taken from the combined DOS of the four DFT calculations in this work and normalized to a per-atom occupation, is presented in Fig. 11. The spin-up and spin-down occupations at the Fermi energy are 0.613 and 0.376 states eV$^{-1}$atom$^{-1}$, respectively, resulting in a combined total Fermi level occupation of $g(E_F) = 1.349$ states eV$^{-1}$atom$^{-1}$. Using this total Fermi level in Eq. (8), one finds $\gamma_0 = 3.18(9)$ mJ mol$^{-1}$ atom K$^{-2}$, leading to an effective carrier mass of $m^* = 4.8(1) m_e$.

This analysis can be extended by first examining the results provided by the Hall studies which provide a carrier density $n = 2.9(8) \times 10^{22}$ cm$^{-3}$. Combined with a value of the effective mass $m^*$ extracted from the specific heat, estimates of the Fermi energy

$$E_F = \frac{\hbar^2}{2m^*}(3\pi^2 n)^{2/3},$$

and the Pauli paramagnetic susceptibility

$$\chi_P = \frac{3\mu_0 \mu_B^2}{2E_F},$$

yield $E_F = 0.72(8)$ eV and $2.3(2) \times 10^{-5}$ emu mol$^{-1}$ atom$^{-1}$, respectively. By comparison, fitting experimental susceptibility data yielded a constant offset of $5.4 \times 10^{-4}$ emu mol$^{-1}$ atom$^{-1}$, as seen in Table II. A diamagnetic contribution to the temperature-independent part of the susceptibility is also expected, consisting of Landau diamagnetic and core diamagnetic components [70]. Landau diamagnetic susceptibility can be determined by

$$\chi_L = -\frac{1}{3} \frac{m_e^2}{\gamma_0} \chi_P,$$

which produces a value of $\chi_L = -3.3(3) \times 10^{-7}$ emu mol$^{-1}$ atom$^{-1}$ [70]. The core diamagnetism depends on the ionization of the atoms within the compound [80], which is currently not known. However, based on the possible ionization states of Cr, Mn, Fe, Co, and Ni, the core diamagnetic susceptibility $\chi_{core}$ must be between $-6.0 \times 10^{-7}$ and $-1.1 \times 10^{-6}$ emu mol$^{-1}$ atom$^{-1}$. These estimates suggest that the effects of Landau diamagnetism are negligible in the overall susceptibility of the equiatomic Cantor alloy, and the core diamagnetism may account for at most a 5% reduction in the temperature-independent susceptibility.

A value for the Pauli paramagnetic susceptibility can also be determined from first-principles methods according to [81]

$$\chi_P = \mu_0 \mu_B g(E_F).$$

This equation gives a Pauli paramagnetic susceptibility of $4.4 \times 10^{-5}$ emu mol$^{-1}$ atom$^{-1}$, similar to the value calculated from Hall data.

A number of factors point toward the possibility of Stoner enhanced paramagnetism as the source of the discrepancy between the calculated Pauli susceptibility and the experimentally determined offset. First, more than half of the component elements of the CrMnFeCoNi alloy are known to be ferromagnetic, and the work of Schneeweiss et al. [26] on this compound identifies the 43 K transition as ferromagnetic, though this claim is disputed by Kamarád et al. [27]. These factors suggest a proximity to magnetic order. Computational results also show a discrepancy in the spin-up and spin-down DOS, which indicates a spontaneous splitting of energy states. Alongside a molecular field of sufficient strength, this splitting will result in a Stoner enhancement [81].

### D. Stoner enhancement of Pauli paramagnetism

The static component of the susceptibility is given by

$$\chi_{static} = \frac{\chi_P}{1 - Z},$$

in which $Z$ is the Stoner enhancement parameter, a value between zero and 1 [81,82]. Assuming that $\chi_P$ from the modified Curie-Weiss fit is an accurate measure of the static component of the susceptibility, the Pauli paramagnetism can be used to calculate $Z$. The $\chi_P$ obtained from Hall data gives a value of $Z = 0.96$. Calculating the Pauli susceptibility from
TABLE IV. Comparison of experimentally determined Stoner enhancement parameters for multiple compounds. The Stoner parameter calculated from Eq. (10) remains at approximately 0.96 regardless of whether the core diamagnetic correction is applied.

| Material                  | \( Z \) | Ref. |
|---------------------------|---------|------|
| CrMnFeCoNi, Eq. (10)     | 0.96    | This work |
| CrMnFeCoNi, Eq. (12)     | 0.92    | This work |
| CrMnFeCoNi, Eq. (15)     | 0.97    | This work |
| Pd                        | 0.82    | [83–86] |
| TiBe2                     | 0.91    | [87]   |
| HfZn2                     | 0.79    | [88]   |
| YFe2Zn20                  | 0.94    | [82]   |
| YCo2Zn20                  | 0.50    | [82]   |
| WB4                       | 0.93    | [86]   |

the computational DOS using Eq. (12) produces a Stoner enhancement of \( Z = 0.92 \). Alternatively, \( Z \) can be calculated using \( \gamma \), the linear component of the specific heat using Eq. (8), in combination with

\[
\chi_p = \mu_0\mu_B^2 g(E_F) \left[ 81 \right], \quad (14)
\]

to obtain an equation for the Stoner enhancement parameter \( Z \) as a function of \( \gamma \), i.e.,

\[
Z = 1 - \frac{3\mu_B^2}{\pi^2 k_B^2} \frac{\gamma}{\chi_{\text{static}}} \left[ 82 \right]. \quad (15)
\]

Using the value of \( \gamma \) obtained from specific heat measurements, this expression gives a value of \( Z = 0.97 \). These Stoner enhancement parameters are listed alongside those of other known compounds listed in Table IV. Additionally, Leong et al. [23] report theoretical values for effective Stoner enhancement parameters for a number of CoCrFeNi-based high-entropy alloys, all of which were found to be either ferromagnetic or close to ferromagnetic ordering.

One may observe a similarity between Eq. (15) and the Wilson ratio

\[
R_W = \frac{4\pi^2 k_B^2}{3\mu_0 g_\mu_B^2} \frac{\chi_0}{\gamma}, \quad (16)
\]

which provides information on the electron-electron correlations in the compound, where a free electron gas has a Wilson ratio of 1 while a higher ratio indicates stronger interactions [89,90]. The experimental results reported here give a Wilson ratio of 2.6 for the Cantor alloy anneal A sample. While this \( R_W \) is higher than the \( R_W \) of 1.2 to 2.2 observed in the strongly correlated electron compound Sr$_2$RuO$_4$, much higher Wilson ratios have been recorded [91]. For example, Balents [92] calculates a value of \( R_W \) as high as 230 in FeSc$_2$S$_3$ based on previously published data [93,94], perhaps due to competition between spin-orbit coupling and magnetic exchange, and proximity of a quantum critical point [92,95]. Similarly, Julian et al. [91] calculate a Wilson ratio of approximately 40 for the nearly ferromagnetic compound Ni$_3$Ga, also using published data [96–98].

E. Magnetic state according to the Stoner criterion

The Stoner model of itinerant magnetism states the ferromagnetic phase should be favored if the Stoner criterion, 

\[ g_0(E_F) I \geq 1, \]

is satisfied, where \( g_0(E_F) \) is the DOS per atom at the Fermi level in the nonmagnetic case (i.e., from a non-spin-polarized calculation). The Stoner exchange parameter \( I \) describes the splitting between the spin-up and spin-down states in the magnetic phase. In ferromagnetic materials, the wavefunctions of the magnetic and nonmagnetic states are identical. However, in the magnetic state, the eigenvalues \( \epsilon \) are shifted by a constant amount \( \pm \frac{1}{2}IM \), where \( M \) is the magnetization [81,99]. Thus the eigenvalues of the up- and down-spin states can be written as

\[
\epsilon_{\uparrow} = \epsilon_0 + \frac{1}{2}IM, \quad \epsilon_{\downarrow} = \epsilon_0 - \frac{1}{2}IM, \quad (17)
\]

where \( k \) and \( v \) denote the wavevectors and band indices. By subtracting Eq. (18) from Eq. (17), one finds \( \Delta = (\epsilon_{\uparrow} - \epsilon_{\downarrow}) = IM \), which can be estimated according to the splitting of the spin-up and down DOS of the magnetic system, measured in eV. Although several methods exist for determining the shift between spin states, there is no exact or designated method, so any Stoner parameter obtained via the DOS should be considered only an approximation. In this work, the overall splitting was measured according to the average shifts in the approximate peak locations within the d-band DOS, yielding \( \Delta \approx 0.233 \text{ eV} \). Combined with a net magnetization of 0.294\( \mu_B \), this \( \Delta \) value provides a Stoner exchange parameter \( I = 0.793 \). To complete the Stoner criterion, the Fermi level occupation of the alloy in a nonmagnetic state is needed. To obtain this value, an additional DFT simulation was performed for the equiatomic alloy in which spin-polarization effects were excluded, yielding \( g_0(E_F) = 1.663 \text{ states/(eV atom)} \) for the nonmagnetic alloy. Evaluation of the Stoner criterion then yields \( g_0(E_F) I = 1.318 \), suggesting that a weakly ferromagnetic phase should be preferred in this system.

F. Magnetic order

The “step” in the susceptibility data visible around 43 K is roughly \( 1 \times 10^{-2} \mu_B \text{ atom}^-1 \) at 0.1 and 1 kOe. This transition is notable in its small size, even relative to transitions in “small-moment” ferromagnets [100–102]. A similar transition was observed in the compound Fe$_{40}$Mn$_{40}$Co$_{10}$Cr$_{10}$ by Egilmez and Abuzaud [69], who identified it as ferrimagnetic and suggested that its small size was due to strong antiferromagnetic coupling between atoms.

Insight about the complex and diverse nature of the local magnetic environments is provided by DFT simulations, performed both in this work and in earlier investigations by others studying very similar alloys [103,104]. Specifically, a sense of the variations of local magnetism can be garnered from Fig. 12, where the local atomic magnetic moments, \( m \), of each atom in our DFT models are plotted against the average local magnetic moment of their first-nearest-neighbor shells, \( m_{NN} \). The atoms are grouped according species type, giving
a picture of the type of coupling (i.e., ferro- or antiferromagnetic) each species prefers to form with its local environment. The total magnetic moment, $\mu_{\text{DFT}}$, from all four simulations, taken as the average moment of all local atomic moments, was found to be 0.294\(\mu_B\), which is much lower than the average magnitude (i.e., disregarding spin direction) of the local moments found to be 1.103\(\mu_B\). This finding, which shows good agreement with previous DFT predictions of moments in the Cantor alloy [105], indicates an uneven level of ferro- and antiferromagnetic couplings within the compound, thereby providing a basis for overall (macroscopic) ferrimagnetic-like ordering.

Examining the couplings exhibited by each species in Fig. 12, one can see that the elements on the left side (i.e., Cr and Mn) of the Periodic Table $d$ block tend to antiferromagnetically align with their local nearest-neighbor shell. In other words, if the spins of an atom’s closest neighbor atoms align on average in one direction, the Cr and Mn atom spins are likely to be found aligned in the opposite direction. Conversely, Co and Ni, residing on the right side of the $d$ block, are more likely to align their spins ferromagnetically, or in the same direction, as the spins of their neighbors. Fe, however, which sits at the center of the $d$ block, is more difficult to classify. The majority of Fe atoms tend to exhibit strongly spin-up configurations, regardless of their immediate local environment. In some rare cases, an Fe atom will take on a strongly spin-down configuration. In all cases, however, the average Weiss field acting on the atoms is in the spin-up direction and as the overwhelming majority of Fe atoms tend to align in this direction, it suggests that Fe may be more sensitive to longer-range magnetic interactions (i.e., beyond the first-nearest-neighbor shell) and that Fe atoms in the Cantor alloy exhibit a more ferromagnetic character. As the magnitude of the Fe-atom moments is so consistently strong, it may be that the noncoupled Fe atoms are what ultimately drive the tilt in spin balance towards one direction, away from an overall antiferromagnetic order and into the ferrimagnetic regime.

Experimentally, the Stoner enhancement and the small magnetic moment suggest that the equiatomic Cantor alloy may be a weak itinerant ferromagnet [106–109]. Furthermore, the estimated magnetic component of entropy calculated from the results in Fig. 10 is approximately 0.096 $R \ln 2$ at 50 K and 0.23 $R \ln 2$ at 97 K, which is an outcome reminiscent of the magnetic entropies of well-known itinerant ferromagnets ZrZn$_2$ and Sc$_3$In (0.02 $R \ln 2$ and 0.005 $R \ln 2$, respectively [110–112]). This small entropy points toward the involvement of a relatively small number of spins indicative of itinerant magnetism. Taken together with evidence of antiferromagnetic coupling from DFT calculations and modified Curie-Weiss fitting to experimental data, these pieces of evidence suggest that the equiatomic Cantor alloy is an itinerant ferrimagnet below 43 K.

V. CONCLUSION

By combining a suite of experimental techniques with numerical simulations, this work extends our understanding of the magnetic properties of the equiatomic Cantor alloy CrMnFeCoNi. Both experimental and DFT results suggest the presence of weak ferrimagnetism below about 43 K, while $\mu$SR measurements indicate a spin-glass-like transition near 85 K. In addition, a large offset in the magnetic susceptibility reveals strong Stoner enhancement of the paramagnetism. Taken with the enhancement of the effective mass and the relatively large Wilson ratio, these results reveal the presence of significant electron-electron interactions within the material.

The transitions observed at 43 and 85 K are highly sensitive to cold working and heat treatment. Although high-temperature anneals reduce the effects of cold working, even 6-day anneals do not completely restore the sample’s original magnetic properties. Despite this sensitivity, measurements on known precipitates demonstrate that these transitions are intrinsic to the compound.

This work emphasizes the need for subsequent investigations along multiple avenues. Most immediately, future work...
might extend Hall experiments to lower temperatures and higher fields and examine specific heat under applied magnetic field. Studies could also be undertaken to investigate the magnetic effects resulting from varying the composition of the Cantor alloy with the goal of controllably tuning the magnetic properties. Lastly, the evidence presented in this work suggests that the distinctive processing dependence of the alloy is not due to the effects of known precipitates. Lattice strain is a possible explanation, but a high-temperature anneal would be expected to eliminate strain effects. Annealing does not restore the original state of the alloy suggesting the presence of a mechanism that will require further research to elucidate.

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[1] B. Cantor, K. B. Kim, and P. J. Warren, Mater. Sci. Forum 386-388, 27 (2002).
[2] J.-W. Yeh, S.-K. Chen, S.-J. Lin, J.-Y. Gan, T.-S. Chin, T.-T. Shun, C.-H. Tsau, and S.-Y. Chang, Adv. Eng. Mater. 6, 299 (2004).
[3] G. S. Firstov, T. A. Kosorukova, Y. N. Koval, and V. V. Odnosum, Mater. Today Proc. 2, S499 (2015).
[4] C.-H. Chen and Y.-J. Chen, Scr. Mater. 162, 185 (2019).
[5] K. Motla, Arushi, P. K. Meena, and R. P. Singh, Semicond. Sci. Technol. 35, 074002 (2022).
[6] Y. Zhang, T. Zuo, Y. Cheng, and P. K. Liaw, Sci. Rep. 3, 1455 (2013).
[7] T. Zuo, M. C. Gao, L. Ouyang, X. Yang, Y. Cheng, R. Feng, S. Chen, P. K. Liaw, J. A. Hawk, and Y. Zhang, Acta Mater. 130, 10 (2017).
[8] B. Cantor, I. T. H. Chang, P. Knight, and A. J. B. Vincent, Mater. Sci. Eng., A 375-377, 213 (2004).
[9] F. Otto, A. Dlouhý, C. Somsen, H. Bei, G. Eggerler, and E. P. George, Acta Mater. 61, 5743 (2013).
[10] M. Laurent-Brocq, A. Akhatova, L. Perriére, S. Chebini, X. Sauvage, E. Leroy, and Y. Champion, Acta Mater. 88, 355 (2015).
[11] M. J. Jang, S. Praveen, H. J. Sung, J. W. Bae, J. Moon, and H. S. Kim, J. Alloys Compd. 730, 242 (2018).
[12] J. H. Kim, K. R. Lim, J. W. Won, Y. S. Na, and H.-S. Kim, Mater. Sci. Eng., A 712, 108 (2018).
[13] S.-W. Kim and J. H. Kim, Mater. Sci. Eng., A 718, 321 (2018).
[14] X. F. Wang, Y. Zhang, F. Qiao, and G. L. Chen, Intermetallics 15, 357 (2007).
[15] K. B. Zhang, Z. Y. Fu, J. Y. Zhang, J. Shi, W. M. Wang, H. Wang, Y. C. Wang, and Q. J. Zhang, J. Alloys Compd. 502, 295 (2010).
[16] Y.-F. Kao, S.-K. Chen, T.-J. Chen, P.-C. Chu, J.-W. Yeh, and S.-I. Lin, J. Alloys Compd. 509, 1607 (2011).
[17] M. S. Lucas, L. Mauger, J. A. Muñoz, Y. Xiao, A. O. Sheets, S. L. Semiatin, J. Horwath, and Z. Turgut, J. Appl. Phys. 109, 07E307 (2011).
[18] S. Singh, N. Wanderka, K. Kiefer, S. Siemensmeyer, and J. Banhart, Ultramicroscopy 111, 619 (2011).
[19] L. Liu, J. B. Zhu, J. C. Li, and Q. Jiang, Adv. Eng. Mater. 14, 919 (2012).
[20] S. G. Ma and Y. Zhang, Mater. Sci. Eng., A 532, 480 (2012).
[21] K. Zhang and Z. Fu, Intermetallics 28, 34 (2012).
[22] M.-H. Tsai, Entropy 15, 5338 (2013).
[23] Z. Leong, J. Z. Wróbel, S. L. Dudarev, R. Goodall, I. Todd, and D. Nguyen-Manh, Sci. Rep. 7, 39803 (2017).
[24] K. Jin, B. C. Sales, G. M. Stocks, G. D. Samolyuk, M. Daene, W. J. Weber, Y. Zhang, and H. Bei, Sci. Rep. 6, 20159 (2016).
[25] P. F. Yu, L. J. Zhang, H. Cheng, H. Zhang, M. Z. Ma, Y. C. Li, G. L. Li, P. K. Liaw, and R. P. Liu, Intermetallics 70, 82 (2016).
[26] O. Schneeweiss, M. Fríák, M. Dudová, D. Holec, M. Šob, D. Kriegner, V. Holý, P. Beran, E. P. George, J. Neugebauer, and A. Dlouhý, Phys. Rev. B 96, 014437 (2017).
[27] J. Kamarád, M. Fríák, J. Kaštíl, O. Schneeweiss, M. Šob, and A. Dlouhý, J. Magn. Magn. Mater. 487, 165333 (2019).
[28] D. Billington, A. D. N. James, E. I. Harris-Lee, D. A. Lagos, D. O’Neill, N. Tsuda, K. Toyoki, Y. Kotani, T. Nakamura, H. Bei, S. Mu, G. D. Samolyuk, G. M. Stocks, J. A. Duffy, J. W. Taylor, S. R. Giblin, and S. B. Dugdale, Phys. Rev. B 102, 174405 (2020).
[29] G. E. Bacon and N. Cowlam, J. Phys. C: Solid State Phys. 2, 238 (1969).
[30] L. Patrick, Phys. Rev. 93, 370 (1954).
[31] CRC Handbook of Chemistry and Physics, 102nd ed., edited by J. R. Rumble (CRC Press, Boca Raton, FL, 2021).
[32] S. J. Blundell, Contemp. Phys. 40, 175 (1999).
[33] A. Suter and B. Wojek, Phys. Proc. 30, 69 (2012).
[34] K. A. Petersen, J. Black, and B. A. Frandsen (unpublished).

[35] A. Yauwanc and P. D. de Réotier, *Muon Spin Rotation, Relaxation, and Resonance: Applications to Condensed Matter*, 1st ed. (Oxford University Press, Oxford, UK, 2011).

[36] G. Kresse and J. Hafner, *Phys. Rev. B* 47, 558 (1993).

[37] G. Kresse and J. Hafner, *Phys. Rev. B* 49, 14251 (1994).

[38] G. Kresse, *J. Non-Cryst. Solids* 192-193, 222 (1995).

[39] P. E. Blochl, *Phys. Rev. B* 50, 17953 (1994).

[40] G. Kresse and D. Joubert, *Phys. Rev. B* 59, 1758 (1999).

[41] J. P. Perdew, K. Burke, and M. Ernzerhof, *Phys. Rev. Lett.* 77, 3865 (1996).

[42] S. H. Wei, L. G. Ferreira, J. E. Bernard, and A. Zunger, *Phys. Rev. B* 42, 9622 (1990).

[43] A. Zunger, S. H. Wei, L. G. Ferreira, and J. E. Bernard, *Phys. Rev. Lett.* 65, 353 (1990).

[44] A. van de Walle, P. Tiwary, M. D. de Jong, D. L. Olson, M. D. Asta, A. Dick, D. Shin, W. Yang, L. Q. Chen, and Z. K. Liu, *Calphad* 42, 13 (2013).

[45] A. Togo and T. Tanaka, *Scr. Mater.* 108, 15 (2015).

[46] E. J. Pickering, R. Muñoz-Moreno, H. J. Stone, and N. G. Jones, *Scr. Mater.* 113, 106 (2016).

[47] F. Otto, A. Dlouhy, K. G. Pradeep, M. Kubenová, D. Raabe, G. Eggeler, and E. P. George, *Acta Mater.* 112, 40 (2016).

[48] F. Otto, N. L. Hanold, and E. P. George, *Intermetallics* 54, 39 (2014).

[49] A. Gali and E. P. George, *Intermetallics* 39, 74 (2013).

[50] M. McElfresh, *Fundamentals of Magnetism and Magnetic Measurements* (Quantum Design, 1994).

[51] B. Schuh, F. Mendez-Martin, B. Völker, E. P. George, H. Clemens, R. Pippan, and A. Hohenwarter, *Acta Mater.* 96, 258 (2015).

[52] Y. J. Li, A. Savan, A. Kostka, H. S. Stein, and A. Ludwig, *Mater. Horiz.* 5, 86 (2018).

[53] Y. Uemura, in *Muon Science: Muons in Physics, Chemistry and Materials*, edited by S. Lee, R. Cywinski, and S. Kilcoyne (Taylor & Francis, New York, 1999).

[54] R. Crook, M. R. Cywinski, *J. Phys.: Condens. Matter* 9, 1149 (1997).

[55] Y. J. Uemura, T. Yamazaki, D. R. Harshman, M. Senba, and R. Crook, *J. Phys.: Condens. Matter* 31-34, 1331 (2018).

[56] G. E. Brodale, R. A. Fisher, W. E. Fogle, N. E. Phillips, and J. van Curen, *J. Magn. Magn. Mater.* 31-34, 1331 (1983).

[57] J. A. Mydosh, *Spin Glasses: An Experimental Introduction* (Taylor & Francis, London, 1993).

[58] A. Tari, *The Specific Heat of Matter at Low Temperatures* (Imperial College Press, London, 2003).

[59] C. Kittel, *Quantum Theory of Solids* (Wiley, New York, 1963).

[60] E. Gopal, *Specific Heats at Low Temperatures* (Plenum Press, New York, 1966).

[61] J. M. D. Coey, S. von Molnar, and R. J. Gambino, *Solid State Commun.* 24, 167 (1977).

[62] J. O. Thomson and J. R. Thompson, *Phys. F: Met. Phys.* 11, 247 (1981).

[63] X. N. Lin, V. A. Bondarenko, G. Cao, and J. W. Brill, *Solid State Commun.* 130, 151 (2004).

[64] V. K. Anand, D. T. Adroja, and A. D. Hillier, *Phys. Rev. B* 85, 014418 (2012).

[65] N. W. Ashcroft and N. D. Mermin, *Solid State Physics* (Holt, Rinehart and Winston, New York, 1976).

[66] J. Singleton, *Band Theory and Electronic Properties of Solids* (Oxford University Press, Oxford, UK, 2001).

[67] M. Egilmez and W. Abuzaid, *Sci. Rep.* 11, 8048 (2021).

[68] L. N. Mulay and E. A. Boudreaux, *Theory and Applications of Molecular Diamagnetism* (Wiley, New York, 1976).

[69] G. Cao, J. E. Crow, R. P. Guertin, P. F. Henning, C. C. Homes, M. Strongin, D. N. Basov, and E. Lochner, *Solid State Commun.* 113, 657 (2000).

[70] H. Kaldarar, E. Royanian, H. Michor, G. Hilshcer, E. Bauer, A. Gribanov, D. Shtepa, P. Rogl, A. Grytsiv, Y. Seropetin, and S. Nesterenko, *Phys. Rev. B* 79, 205104 (2009).

[71] T. Grofi, A. W. Pacyna, and E. Malicka, *Solid State Phenom.* 170, 213 (2011).

[72] M. M. Markina, B. V. Mill, E. A. Zvereva, A. V. Usakhow, S. V. Streltsov, and A. N. Vasiliev, *Phys. Rev. B* 89, 104409 (2014).

[73] D. V. Schroeder, *An Introduction to Thermal Physics*, Int. ed. (Addison-Wesley, San Francisco, CA, 2000).

[74] C. Kittel, *Introduction to Solid State Physics*, 8th ed. (Wiley, New York, 2005).

[75] J. J. Sabo, *Phys. Rev. B* 1, 1325 (1970).

[76] J. M. Ziman, *Principles of the Theory of Solids*, 2nd ed. (University Press, Cambridge, UK, 1972).

[77] B. Hammer and J. K. Norskog, *Nature (London)* 376, 238 (1995).

[78] G. A. Bain and J. F. Berry, *J. Chem. Educ.* 53, 532 (2008).

[79] S. Blundell, *Magnetism in Condensed Matter* (Oxford University Press, Oxford, UK, 2001).

[80] S. Jia, S. L. Bud’ko, G. D. Samolyuk, and P. C. Canfield, *Nat. Phys.* 3, 334 (2007).

[81] F. E. Hoare, J. C. Matthews, J. C. Walling, and E. C. Stoner, *Proc. R. Soc. London A* 216, 502 (1953).

[82] G. Chouteau, R. Fourneaux, K. Gobrecht, and R. Tournier, *Phys. Rev. Lett.* 20, 193 (1968).

[83] S. C. Hong, T. H. Rho, and J. Il Lee, *J. Magn. Magn. Mater.* 140-144, 697 (1995).

[84] J. Shen, J. Zhou, M. Zhao, G. Liu, J. Gao, and J. Yao, *J. Phys.: Condens. Matter* 28, 026005 (2016).

[85] G. R. Stewart, J. L. Smith, A. L. Giorgi, and Z. Fisk, *Phys. Rev. B* 25, 5907 (1982).

[86] H. B. Radousky, G. S. Knapp, J. W. Downey, A. T. Aldred, and A. J. Freeman, *J. Magn. Magn. Mater.* 40, 117 (1983).

[87] J. Sólyom, *Fundamentals of the Physics of Solids* (Springer, Berlin, 2009).

[88] U. B. Paramanik, Anupam, U. Burkhardt, R. Prasad, C. Geibel, and Z. Hossain, *J. Alloys Compd.* 580, 435 (2013).

[89] S. R. Julian, A. P. Mackenzie, G. G. Lonzarich, C. Bergemann, R. K. W. Haselwimmer, Y. Maeno, S. NishiZaki, A. W. Tyler,
