



Effective One-Dimensional Coupling in the Highly Frustrated Square-Lattice Itinerant Magnet $\text{CaCo}_{2-y}\text{As}_2$

A. Sapkota,^{1,2} B. G. Ueland,^{1,2} V. K. Anand,^{1,3} N. S. Sangeetha,^{1,2} D. L. Abernathy,⁴ M. B. Stone,⁴ J. L. Niedziela,⁵ D. C. Johnston,^{1,2} A. Kreyssig,^{1,2} A. I. Goldman,^{1,2} and R. J. McQueeney^{1,2}

¹Ames Laboratory, U.S. Department of Energy, Iowa State University, Ames, Iowa 50011, USA

²Department of Physics and Astronomy, Iowa State University, Ames, Iowa 50011, USA

³Helmholtz-Zentrum Berlin für Materialien und Energie GmbH, Hahn-Meitner Platz 1, D-14109 Berlin, Germany

⁴Quantum Condensed Matter Division, Oak Ridge National Laboratory, Oak Ridge, Tennessee 37831, USA

⁵Instrument and Source Division, Oak Ridge National Laboratory, Oak Ridge, Tennessee 37831, USA

(Received 17 January 2017; published 2 October 2017)

Inelastic neutron scattering measurements on the itinerant antiferromagnet $\text{CaCo}_{2-y}\text{As}_2$ at a temperature of 8 K reveal two orthogonal planes of scattering perpendicular to the Co square lattice in reciprocal space, demonstrating the presence of effective one-dimensional spin interactions. These results are shown to arise from near-perfect bond frustration within the J_1 - J_2 Heisenberg model on a square lattice with ferromagnetic J_1 and hence indicate that the extensive previous experimental and theoretical study of the J_1 - J_2 Heisenberg model on local-moment square spin lattices should be expanded to include itinerant spin systems.

DOI: 10.1103/PhysRevLett.119.147201

Magnetic frustration arises when competing interactions between magnetic moments (spins) cannot be mutually satisfied. It suppresses the development of long-range magnetic order and often creates enhanced spin fluctuations, which can lead to a variety of novel phases including quantum spin liquids [1,2], spin and electronic nematic phases, and unconventional superconductivity [3–6]. There are many examples of materials in which geometry of the lattice leads to frustration, such as pyrochlore, spinel, or kagome systems [1,7]. However, in the case of a square-lattice system, the frustration can arise from competing nearest-neighbor (NN) and next-nearest-neighbor (NNN) interactions [8,9].

Compounds with the chemical formula ATM_2As_2 (with $A = \text{Ca, Sr, Ba}$ and $\text{TM} = \text{Mn, Fe, Co, Ni}$) form a large class of quasi-two-dimensional (quasi-2D) materials containing layers of TM ions on a square lattice, which are stacked along c . Despite the crystal structure being three dimensional, they are considered quasi-2D for magnetism, as the interactions between layers are much smaller than those within the layers. Much of the recent motivation for the study of these materials is due to the proximity of antiferromagnetic (AFM) order and high-temperature superconductivity in the doped variants of $\text{TM} = \text{Fe}$ compounds [10–13]. The ATM_2As_2 materials adopt several different magnetic structures, including Néel-(or checkerboard)type AFM (e.g., BaMn_2As_2 [14]), stripe-type AFM (e.g., AFe_2As_2 [13]), and A-type AFM (e.g., $\text{CaCo}_{2-y}\text{As}_2$ [15]). The AFM order in the $\text{TM} = \text{Fe}$ and Co variants is itinerant in nature, possessing an ordered moment of $\mu \lesssim 1\mu_B/\text{Fe}$ [13] and $< 0.6\mu_B/\text{Co}$ [15], respectively. The itinerancy of $\text{CaCo}_{2-y}\text{As}_2$ is inferred from the resistivity,

magnetization, angle-resolved photoemission spectroscopy measurements and band structure calculations [16].

Despite their itinerant nature, the magnetic structures and spin fluctuations can be minimally described by considering NN (J_1) and NNN (J_2) [17] magnetic exchange interactions between magnetic ions on a square lattice [18–20]. In general, the magnetic ground state is determined by the relative strengths of J_1 and J_2 , with Néel-type, stripe-type AFM, and ferromagnetic (FM)/A-type ordering occurring for $J_1 > 2J_2$ ($J_1 > 0$), $|J_1| < 2J_2$ ($J_2 > 0$), and $-J_1 > 2J_2$ ($J_1 < 0$), respectively [8,13]. The system becomes frustrated and ordering in any of these magnetic structures is suppressed when $|J_1| \approx 2J_2$ and $J_2 > 0$ (AFM). A frustration parameter, $\eta = J_1/2J_2$, quantifies the level of the magnetic frustration. Maximum frustration occurs when $\eta = 1$ (–1), and the stripe-type and Néel-type (stripe-type and FM) ground states compete [8,13].

The frustration parameter η can be determined experimentally by measuring the spatial anisotropy of the spin fluctuations using inelastic neutron scattering (INS). The spatial distributions of the spin fluctuations in different magnetic ground states depend strongly on η as illustrated in the Supplemental Material [21] (Fig. S4). For example, INS measurements for the parent and doped compositions of AFe_2As_2 compounds find η in the range from 0.3–0.7 [22,23], suggesting the presence of a moderate degree of frustration. INS experiments on SrCo_2As_2 observe stripe-type AFM spin fluctuations peaked at $\mathbf{Q}_{\text{stripe}}$ similar to AFe_2As_2 [24]. The anisotropy of the spin fluctuations gives $\eta \approx -0.6$ [24] and indicates moderate frustration.

Here, we report inelastic neutron scattering measurements of magnetic excitations in $\text{CaCo}_{2-y}\text{As}_2$ at $T = 8$ K,

below the AFM ordering temperature $T_N = 52$ K [15,16], that reveal *planes* of magnetic scattering in reciprocal space perpendicular to the Co square-lattice planes. This is in sharp contrast to *rods* of scattering expected perpendicular to quasi-two-dimensional planes of spins but similar to the expectation for quasi-one-dimensional spin chains. From this anisotropy, we model the data and estimate $\eta \approx -1$, corresponding to extreme frustration. Itinerant $\text{CaCo}_{2-y}\text{As}_2$ stands unique among the other square-lattice systems as the maximum frustration parameter observed in the most-well-studied insulating local-moment oxides is -0.56 [25], far from perfect frustration. These observations indicate that previous extensive studies of the J_1 - J_2 Heisenberg model on local-moment square spin lattices should be expanded beyond oxides to include itinerant spin systems found in arsenides and potentially other pnictides. This has many potentially interesting ramifications and applications.

Single crystals of $\text{CaCo}_{2-y}\text{As}_2$ ($y \approx 0.14$) were grown using Sn flux [15,16]. The vacancies on the Co sites are randomly distributed, as x-ray and neutron diffraction measurements find no evidence for vacancy ordering [15,16]. INS experiments were performed on the ARCS spectrometer [26] at the Spallation Neutron Source, at Oak Ridge National Laboratory. Six single crystals of

$\text{CaCo}_{2-y}\text{As}_2$ with a total mass of ~ 1.1 g were coaligned in the $(H, 0, L)$ scattering plane (in trigonal notation) with a full width at half maximum of less than 4° . The coaligned set was attached to the cold finger of a closed-cycle He cryostat. The measurements were carried out with the incident beam along the c axis and incident energies of $E_i = 75$ and 250 meV at $T = 8$ K. Throughout the Letter, we define $\mathbf{Q} = (H - K, H + K, L) = (2\pi/a)(H - K)\hat{\mathbf{i}} + (2\pi/a)(H + K)\hat{\mathbf{j}} + (2\pi/c)L\hat{\mathbf{k}}$ with respect to the tetragonal $I4/mmm$ crystal system, where $a = 3.98$ Å and $c = 10.27$ Å. In this notation, the reciprocal lattice vectors for stripe and A-type magnetic order are $\mathbf{Q}_{\text{stripe}} = (\frac{1}{2}, \frac{1}{2}, 1) + (m, n, l)$ with m, n integers and l even and $\mathbf{Q}_A = (H - K, H + K, L)$ where $H - K$ and $H + K$ are integers with $|H - K| + |H + K|$ even and L odd. The data were visualized using the MSLICE software [27].

Figures 1(a)–1(c), 1(g), and 1(h) are the INS data measured on ARCS and show the presence of very striking spin fluctuations in $\text{CaCo}_{2-y}\text{As}_2$. $\text{CaCo}_{2-y}\text{As}_2$ has A-type AFM order at low temperatures and the low-energy spin fluctuations are expected to originate at $\mathbf{Q}_A = [(0, 0), (1, 1)]$, using 2D notation. However, constant-energy slices of the data in Figs. 1(a)–1(c) show that the scattering from the spin fluctuations does not occur only

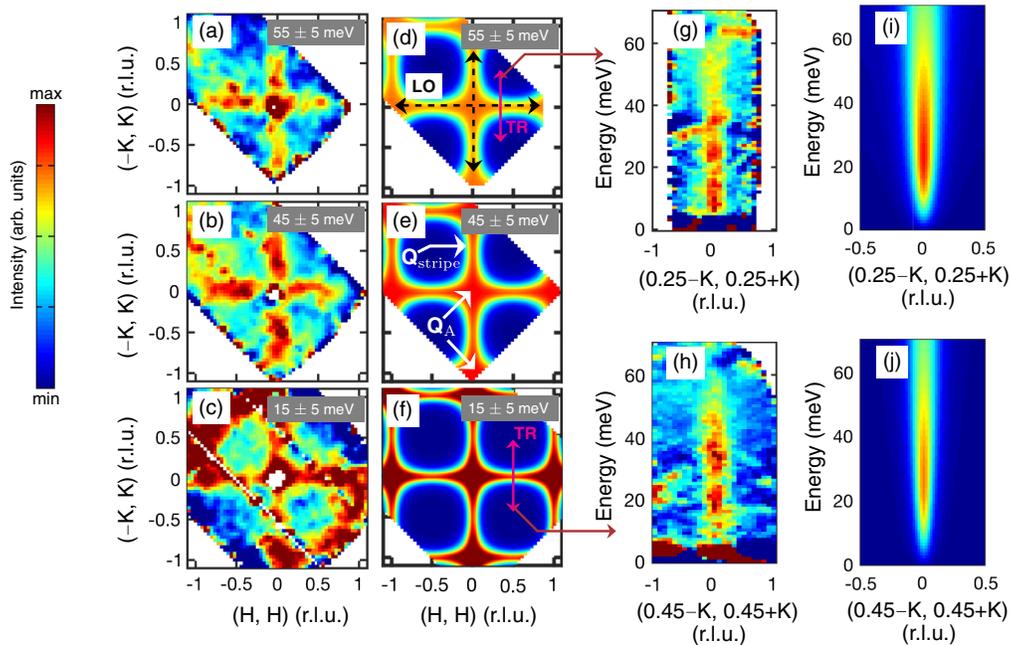


FIG. 1. Spin fluctuations in $\text{CaCo}_{2-y}\text{As}_2$: Experimental INS data ($E_i = 75$ meV) vs diffusive model. (a)–(c) Constant-energy slices of the background-subtracted data (the isotropic background is estimated similar to Ref. [22]) averaged over the energy ranges of 50–60, 40–50, and 10–20 meV, respectively. (d)–(f) Corresponding constant-energy slices calculated using the diffusive model and parameters obtained from fitting the data shown in Fig. 2. In (d), the black dashed arrows show the LO directions and the maroon arrow shows the TR direction. (g),(h) Transverse slices of the background-subtracted data along the $[-K, K]$ direction through $(0.25, 0.25)$ and $(0.45, 0.45)$ (r.l.u.), respectively, obtained after averaging over the LO direction by ± 0.1 r.l.u. (i),(j) Corresponding energy dependence of the scattering obtained using the diffusive model and parameters of fits to the data in Fig. 2. The energy-dependent figures are averaged over all four symmetry-related quadrants. The color scale represents the intensity in each panel. The maximum/minimum intensities in the panels are (a) 0.34/ -0.5 , (b) 0.42/ -0.3 , (c) 0.65/ -0.15 , (d)–(f) 0.43/0, (g) 0.52/ -0.45 , (h) 0.65/ -0.35 , and (i),(j) 0.52/0 in arbitrary units.

around positions corresponding to \mathbf{Q}_A , but extends along the longitudinal (LO) direction, crossing positions including $\mathbf{Q}_{\text{stripe}} = (1/2, 1/2)$, at all accessible energies from ~ 10 – 120 meV. In contrast, the scattering along the transverse (TR) direction is sharp. The energy dependence [Figs. 2(g) and 2(h) and Supplemental Material [21] Fig. S5] shows steep dispersion in the TR direction and the scattering extends in energy beyond 100 meV. As we will show below, the extreme anisotropy of the spin fluctuations gives $\eta \approx -1$, implying that $\text{CaCo}_{2-y}\text{As}_2$ exhibits near-perfect magnetic frustration between FM/A-type and stripe-type ordering.

To describe the scattering data, we consider two models for A-type AFM. Using values of J_1 and J_2 appropriate for an A-type AFM ground state, we first use the linear spin-wave theory approximation to the Heisenberg model to calculate the neutron scattering cross section corresponding to the values of \mathbf{Q} and E measured by INS. The details and results of the model calculations are shown in the Supplemental Material [21] Fig. S1. We find that when adopting parameters corresponding to nearly maximal

frustration ($\eta \approx -1$), the spin waves collapse along the LO direction, leading to a spin-wave anisotropy that is similar to the experimental data at low energies. However, the calculated cross section and INS data show significant differences at higher energies (Supplemental Material [21] Fig. S1).

We next consider a model more appropriate for itinerant systems close to magnetic order. In itinerant magnets, the electronic degrees of freedom can result in a significant degree of Landau damping of the spin fluctuations due to the scattering of electrons. A diffusive model that describes such overdamped spin fluctuations in a nearly ordered system has been used to describe INS data on weakly ordered and metallic $\text{Ba}(\text{Fe}_{1-x}\text{Co}_x)_2\text{As}_2$ [22,28,29] and paramagnetic CaFe_2As_2 [30]. Neglecting the weak AFM interlayer interactions and keeping intra-layer interactions up to NNN only, we develop a similar model for FM/A-type fluctuations. We find that the imaginary part of the generalized magnetic susceptibility, which is proportional to the INS spectrum, can be written as

$$\chi''(\mathbf{q}, \omega) = \frac{\chi_0 \Gamma \omega}{\Gamma^2 (1 + \xi_q^2 q^2)^2 + \omega^2}, \quad (1)$$

where

$$\xi_q^2 q^2 = (2\xi/a)^2 \left\{ \eta \left[\cos\left(\frac{q_x + q_y}{2} a\right) + \cos\left(\frac{q_y - q_x}{2} a\right) \right] + \cos\left(\frac{q_x + q_y}{2} a\right) \cos\left(\frac{q_y - q_x}{2} a\right) - 2\eta - 1 \right\}. \quad (2)$$

Here, χ_0 is the static uniform susceptibility, ξ is the magnetic correlation length, Γ is the Landau damping parameter, and the x and y directions correspond to the $[1 \ 0 \ 0]$ and $[0 \ 1 \ 0]$ directions of the tetragonal $I4/mmm$ crystal system, respectively. In the itinerant picture relevant for the iron arsenides, η arises from the spatial anisotropy of electronic velocities at the Fermi surface [22,31].

Similar to Ref. [30], we can experimentally determine $\eta = (\xi_{\text{LO}}^2 + \xi_{\text{TR}}^2 / \xi_{\text{LO}}^2 - \xi_{\text{TR}}^2)$ in terms of the magnetic correlation lengths in the LO and TR directions. The magnetic correlation length $\xi_{\text{LO/TR}}$ is inversely proportional to the width of the peak along the respective directions. The almost flat peaks in Figs. 2(a) and 2(c) along the LO direction suggest $\xi_{\text{LO}} \approx 0$, thus indicating $\eta \approx -1$. Moreover, the values of η and other parameters in Eqs. (1) and (2) were determined by fitting the cuts from the data as shown in Fig. 2 (details in the Supplemental Material [21]). The data are best fit with a constant (\mathbf{Q} -independent) damping parameter [$\Gamma = 20(4)$ meV] and $\eta = -1.03(2)$. This value of η indicates the presence of extreme frustration in $\text{CaCo}_{2-y}\text{As}_2$. The calculated neutron scattering cross sections, using the diffusive model, capture all of the essential features corresponding to the INS data shown in Figs. 1 and 2.

Also from the dispersion of A-type AFM in the TR direction, we estimate that $SJ_1 = -52 \pm 3$ meV, where S is the magnitude of the spin [details of estimation are in the Supplemental Material [21], Eqs. (20) and (21)]. We find that the energy scale for the A-type AFM order is much smaller than the FM NN exchange energy ($k_B T_N / |SJ_1| \approx 0.1$). This ratio, 0.1, is much smaller than the same ratio of the moderately frustrated square-lattice oxides (minimum value in oxides is 0.6) [25] and the iron arsenides (0.7) [32]. Thus $\text{CaCo}_{2-y}\text{As}_2$ orders at a temperature much lower than expected based on the strength of its magnetic interactions. This is a hallmark of strong magnetic frustration.

Materials such as $\text{CaCo}_{2-y}\text{As}_2$ are considered quasi-2D because interactions between Co layers are much smaller than those within the layer. In the case of $\text{CaCo}_{2-y}\text{As}_2$, frustrated interactions within the layer reduce the dimensionality even further, leading to effectively 1D behavior. This is most easily pictured by considering J_1 and J_2 interactions in a single square layer with the stripe-type AFM order, as shown in Fig. 3(a). The stripe-type magnetic structure consists of alternating chains of FM spins oriented in the TR direction [ordering vector $(1/2, 1/2)$] with an effective interchain coupling of $\sim |J_1| - 2J_2$ which goes to zero as $\eta \rightarrow (-1)^+$. Moreover, the cost of flipping one FM

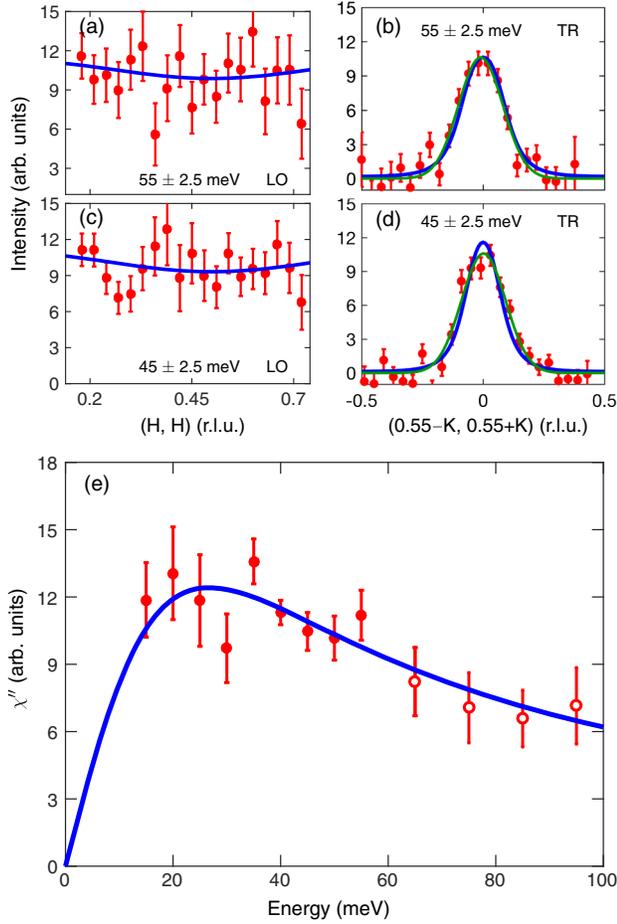


FIG. 2. Constant-energy cuts and dynamical susceptibility of the data fitted with the diffusive model. (a),(c) LO and (b),(d) TR cuts of data through (0.55, 0.55) at (a),(b) 52.5–57.5 and (c),(d) 42.5–47.5 meV. TR cuts were averaged from 0.3–0.8 r.l.u. in the LO direction and LO cuts ± 0.1 r.l.u. in the TR directions. Both data are corrected for the Co^{+1} magnetic form factor after subtracting a background contribution estimated from Gaussian fits to the TR cuts. The single peaked TR cuts are an outcome of the unresolvable broader dispersion branches due to the diffusive nature of the spin fluctuations. The green lines in the TR cuts are fits to a Gaussian line shape and blue solid lines are fits to the diffusive model. (e) Energy dependence of the imaginary part of the dynamical susceptibility centered at (0.55, 0.55) and averaged along the LO direction from 0.3–0.8 r.l.u. Each data point is obtained after fitting the TR cuts with a Gaussian line shape. The closed symbols are data measured with $E_i = 75$ meV and open symbols with $E_i = 250$ meV.

chain against another goes to zero and the FM chains decouple. This results in a vanishing dispersion in the LO direction, but the steep dispersion remains in the TR direction as shown in Fig. 3(c) where effective coupling is maximized as $\sim -|J_1| - 2J_2$. Rodlike fluctuations occur as a result of the effective zero coupling along the LO direction as shown in Fig. 3(c).

Starting from the FM side, the spin stiffness goes to zero in all directions as $\eta \rightarrow (-1)^-$, preserving the fourfold

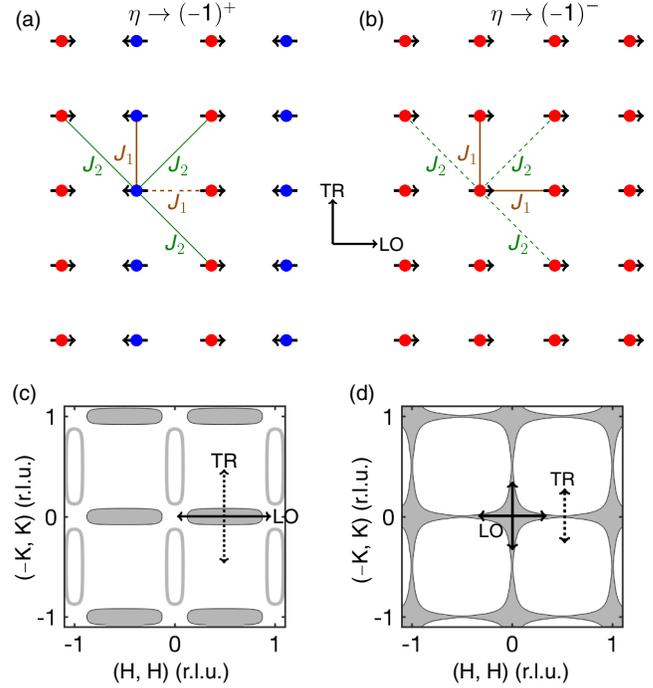


FIG. 3. (a),(b) Coupling of spins on a square lattice corresponding to stripe-type AFM and FM/A-type spin dynamics, respectively, for an extreme degree of frustration showing effective zero coupling along the LO directions. Red and blue circles represent transition metals on different magnetic sublattices. Magnetic NN (J_1) and NNN (J_2) interactions are shown where J_1 is FM and J_2 is AFM. Stable (frustrated) bonds are denoted by a solid (dashed) lines. (c),(d) Spin fluctuations corresponding to highly frustrated spin interactions in (a) and (b), respectively. The empty rods shown in (c) are due to the twin domain of (a), that has alternating chains of blue and red circles along the perpendicular directions. Both occur with equal populations.

symmetry of the ground state. However, similar to the stripe-type AFM order, the effective coupling between FM chains in the LO direction goes to zero, as shown in Fig. 3(b), and any TR component is steeply dispersive. Thus, the low-energy magnetic spectral weight is confined to a wall along the LO direction [Fig. 3(d)], the scattering signature of a 1D system.

In summary, we show that $\text{CaCo}_{2-y}\text{As}_2$ possesses spin fluctuations that are unique compared to AFe_2As_2 and SrCo_2As_2 in that it displays extreme spatial anisotropy. This extreme spatial anisotropy is due to the perfect magnetic frustration arising from the competing FM and AFM interactions and leads to effectively 1D behavior. Also, the value of the frustration parameter $\eta = -1.03(2)$ is in the region where the possibility of spin liquids is discussed [8]. Perfect magnetic frustration, spin liquids, and the exotic properties related to them are extensively discussed and realized in strongly correlated (local-moment) systems. However, the role of magnetic

frustration in moderately/weakly correlated metals is poorly understood. In this regime, coupling of charge carriers to quantum spin fluctuations can lead to many interesting quantum phenomena, including unconventional superconductivity. $\text{CaCo}_{2-y}\text{As}_2$ is a very unusual metallic square-lattice compound in which nearly perfect frustration occurs. The challenge is to identify other potential candidates. For example, one could dope insulators or semiconductors known to be near maximal frustration into the metallic state to generate materials that retain the frustration and in which novel electronic and/or magnetic phenomena may be found.

We are grateful for assistance of G. S. Tucker (Paul Scherrer Institute), Abhishek Pandey (Texas A&M University), Yongbin Lee (Ames Laboratory), Jong Keum (X-ray Laboratory, SNS, ORNL), Songxue Chi (TAX, HFIR, ORNL), and Rafael M. Fernandes (University of Minnesota). Work at the Ames Laboratory was supported by the U.S. Department of Energy, Basic Energy Sciences, Division of Materials Sciences and Engineering, under Contract No. DE-AC02-07CH11358. This research used resources at the Spallation Neutron Source, a DOE Office of Science User Facility operated by the Oak Ridge National Laboratory.

-
- [1] L. Balents, *Nature (London)* **464**, 199 (2010).
- [2] L. Savary and L. Balents, *Rep. Prog. Phys.* **80**, 016502 (2017).
- [3] P. Chandra, P. Coleman, and A. I. Larkin, *Phys. Rev. Lett.* **64**, 88 (1990).
- [4] C. Xu, M. Müller, and S. Sachdev, *Phys. Rev. B* **78**, 020501 (2008).
- [5] C. Fang, H. Yao, W.-F. Tsai, J. P. Hu, and S. A. Kivelson, *Phys. Rev. B* **77**, 224509 (2008).
- [6] N. Shannon, T. Momoi, and P. Sindzingre, *Phys. Rev. Lett.* **96**, 027213 (2006).
- [7] J. E. Greedan, *J. Mater. Chem.* **11**, 37 (2001).
- [8] N. Shannon, B. Schmidt, K. Penc, and P. Thalmeier, *Eur. Phys. J. B* **38**, 599 (2004).
- [9] R. Nath, Y. Furukawa, F. Borsa, E. E. Kaul, M. Baenitz, C. Geibel, and D. C. Johnston, *Phys. Rev. B* **80**, 214430 (2009).
- [10] A. S. Sefat, R. Jin, M. A. McGuire, B. C. Sales, D. J. Singh, and D. Mandrus, *Phys. Rev. Lett.* **101**, 117004 (2008).
- [11] A. Leithe-Jasper, W. Schnelle, C. Geibel, and H. Rosner, *Phys. Rev. Lett.* **101**, 207004 (2008).
- [12] N. Kumar, R. Nagalakshmi, R. Kulkarni, P. L. Paulose, A. K. Nigam, S. K. Dhar, and A. Thamizhavel, *Phys. Rev. B* **79**, 012504 (2009).
- [13] D. C. Johnston, *Adv. Phys.* **59**, 803 (2010).
- [14] D. C. Johnston, R. J. McQueeney, B. Lake, A. Honecker, M. E. Zhitomirsky, R. Nath, Y. Furukawa, V. P. Antropov, and Y. Singh, *Phys. Rev. B* **84**, 094445 (2011).
- [15] D. G. Quirinale, V. K. Anand, M. G. Kim, A. Pandey, A. Huq, P. W. Stephens, T. W. Heitmann, A. Kreyssig, R. J. McQueeney, D. C. Johnston, and A. I. Goldman, *Phys. Rev. B* **88**, 174420 (2013).
- [16] V. K. Anand, R. S. Dhaka, Y. Lee, B. N. Harmon, A. Kaminski, and D. C. Johnston, *Phys. Rev. B* **89**, 214409 (2014).
- [17] Here and we define $J < 0$ for FM and $J > 0$ for AFM exchange interactions.
- [18] R. J. McQueeney, S. O. Diallo, V. P. Antropov, G. D. Samolyuk, C. Broholm, N. Ni, S. Nandi, M. Yethiraj, J. L. Zarestky, J. J. Pulikotil, A. Kreyssig, M. D. Lumsden, B. N. Harmon, P. C. Canfield, and A. I. Goldman, *Phys. Rev. Lett.* **101**, 227205 (2008).
- [19] J. Zhao, D. T. Adroja, D.-X. Yao, R. Bewley, S. Li, X. F. Wang, G. Wu, X. H. Chen, J. Hu, and P. Dai, *Nat. Phys.* **5**, 555 (2009).
- [20] E. Manousakis, J. Ren, S. Meng, and E. Kaxiras, *Solid State Commun.* **150**, 62 (2010).
- [21] See Supplemental Material at <http://link.aps.org/supplemental/10.1103/PhysRevLett.119.147201> for details about the Heisenberg model, diffusive model and estimation of SJ_1 and fitting of the INS data.
- [22] G. S. Tucker, R. M. Fernandes, H.-F. Li, V. Thampy, N. Ni, D. L. Abernathy, S. L. Bud'ko, P. C. Canfield, D. Vaknin, J. Schmalian, and R. J. McQueeney, *Phys. Rev. B* **86**, 024505 (2012).
- [23] L. W. Harriger, H. Q. Luo, M. S. Liu, C. Frost, J. P. Hu, M. R. Norman, and P. Dai, *Phys. Rev. B* **84**, 054544 (2011).
- [24] W. Jayasekara, Y. Lee, A. Pandey, G. S. Tucker, A. Sapkota, J. Lamsal, S. Calder, D. L. Abernathy, J. L. Niedziela, B. N. Harmon, A. Kreyssig, D. Vaknin, D. C. Johnston, A. I. Goldman, and R. J. McQueeney, *Phys. Rev. Lett.* **111**, 157001 (2013).
- [25] R. Nath, A. A. Tsirlin, H. Rosner, and C. Geibel, *Phys. Rev. B* **78**, 064422 (2008).
- [26] D. L. Abernathy, M. B. Stone, M. J. Loguillo, M. S. Lucas, O. Delaire, X. Tang, J. Y. Y. Lin, and B. Fultz, *Rev. Sci. Instrum.* **83**, 015114 (2012).
- [27] Developed by Radu Coldea 1998–2001 and ISIS add ons and enhancements 2001–2014, <http://mslice.isis.rl.ac.uk/>.
- [28] D. S. Inosov, J. T. Park, P. Bourges, D. L. Sun, Y. Sidis, A. Schneidewind, K. Hradil, D. Haug, C. T. Lin, B. Keimer, and V. Hinkov, *Nat. Phys.* **6**, 178 (2010).
- [29] G. S. Tucker, R. M. Fernandes, D. K. Pratt, A. Thaler, N. Ni, K. Marty, A. D. Christianson, M. D. Lumsden, B. C. Sales, A. S. Sefat, S. L. Bud'ko, P. C. Canfield, A. Kreyssig, A. I. Goldman, and R. J. McQueeney, *Phys. Rev. B* **89**, 180503 (2014).
- [30] S. O. Diallo, D. K. Pratt, R. M. Fernandes, W. Tian, J. L. Zarestky, M. Lumsden, T. G. Perring, C. L. Broholm, N. Ni, S. L. Bud'ko, P. C. Canfield, H.-F. Li, D. Vaknin, A. Kreyssig, A. I. Goldman, and R. J. McQueeney, *Phys. Rev. B* **81**, 214407 (2010).
- [31] J. Knolle, I. Eremin, A. V. Chubukov, and R. Moessner, *Phys. Rev. B* **81**, 140506 (2010).
- [32] S. O. Diallo, V. P. Antropov, T. G. Perring, C. Broholm, J. J. Pulikotil, N. Ni, S. L. Bud'ko, P. C. Canfield, A. Kreyssig, A. I. Goldman, and R. J. McQueeney, *Phys. Rev. Lett.* **102**, 187206 (2009).