

Magnetic frustration in iridium spinel compound CuIr_2S_4

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Geometrical frustration in electronic degrees of freedom has been one of the major topics in the field of condensed matter physics, for which inorganic compounds with the AB_2X_4 cubic spinel structure consisting of corner-shared network of transition metal ions (A and/or B) have served as fascinating stages. The thiospinel compound, CuIr_2S_4 , is such a recent example, in which a charge order of mixed-valent Ir ions into isomorphous octamers of $\text{Ir}_8^{3+}\text{S}_{24}$ and $\text{Ir}_8^{4+}\text{S}_{24}$ is realized upon metal-insulator (MI) transition at $T_{\text{MI}} = 230$ K[1]. As per the currently accepted scenario regarding the t_{2g} manifold, the frustration is relieved by the formation of Ir^{4+} ($5d^5$, $S = 1/2$) dimers that accompany the spin-singlet ground state driven by orbital order and the associated spin-Peierls instability[2].

Here, we demonstrate via a muon spin rotation and NMR experiments that the electronic ground state of CuIr_2S_4 is not the presumed spin-singlet state but a novel paramagnetic state that exhibits a quasistatic spin glass-like magnetism below ~ 100 K (see Fig.1)[3]. The present result indicates that the geometrical frustration remains partially unresolved even after Ir^{4+} dimarization, suggesting that strong spin-orbit interaction may be playing an important role in determining the ground state.

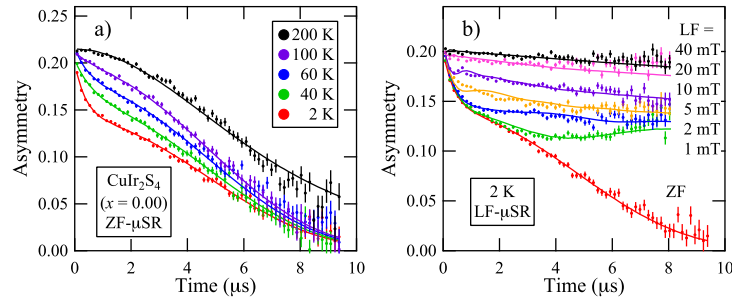


FIG. 1: a) μSR spectra in CuIr_2S_4 under zero external field, observed at several temperatures. b) μSR spectra at 2 K under a longitudinal field up to 40 mT.

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