Novel Magnetism of Ir$^{5+}(5d^4)$ Ions in the Double Perovskite Sr$_2$YIrO$_6$

Gang Cao  
*University of Kentucky*, cao@pa.uky.edu

Tongfei Qi  
*University of Kentucky*, tongfei.qi@uky.edu

Li Li  
*University of Kentucky*, li.li@uky.edu

Jsaminka Terzic  
*University of Kentucky*, jasminka.terzic@uky.edu

Shujuan Yuan  
*University of Kentucky*, syu224@uky.edu

See next page for additional authors  
Follow this and additional works at: [https://uknowledge.uky.edu/physastron_facpub](https://uknowledge.uky.edu/physastron_facpub)

Part of the Astrophysics and Astronomy Commons, and the Physics Commons  
Right click to open a feedback form in a new tab to let us know how this document benefits you.

Repository Citation  
Cao, Gang; Qi, Tongfei; Li, Li; Terzic, Jsaminka; Yuan, Shujuan; De Long, Lance E.; Murthy, Ganpathy; and Kaul, Ribhu K., "Novel Magnetism of Ir$^{5+}(5d^4)$ Ions in the Double Perovskite Sr$_2$YIrO$_6$" (2014). *Physics and Astronomy Faculty Publications*. 3.  
[https://uknowledge.uky.edu/physastron_facpub/3](https://uknowledge.uky.edu/physastron_facpub/3)

This Article is brought to you for free and open access by the Physics and Astronomy at UKnowledge. It has been accepted for inclusion in Physics and Astronomy Faculty Publications by an authorized administrator of UKnowledge. For more information, please contact UKnowledge@lsv.uky.edu.
Novel Magnetism of Ir$^{5+}(5d^4)$ Ions in the Double Perovskite Sr$_2$YIrO$_6$

Digital Object Identifier (DOI)
http://dx.doi.org/10.1103/PhysRevLett.112.056402

Notes/Citation Information
Published in Physical Review Letters, v. 112, no. 5, 056402.

© 2014 American Physical Society.

The copyright holder has granted the permission for posting the article here.

Authors
Gang Cao, Tongfei Qi, Li Li, Jsaminka Terzic, Shujuan Yuan, Lance E. De Long, Ganpathy Murthy, and Ribhu K. Kaul

This article is available at UKnowledge: https://uknowledge.uky.edu/physastron_facpub/3
Novel Magnetism of Ir$^{5+}(5d^4)$ Ions in the Double Perovskite Sr$_2$YIrO$_6$


Department of Physics and Astronomy and Center for Advanced Materials, University of Kentucky, Lexington, Kentucky 40506, USA

Department of Physics, Shanghai University, Shanghai, China

(Received 8 November 2013; published 6 February 2014)

We synthesize and study single crystals of a new double-perovskite Sr$_2$YIrO$_6$. Despite two strongly unfavorable conditions for magnetic order, namely, pentavalent Ir$^{5+}(5d^4)$ ions which are anticipated to have $J_{\text{eff}} = 0$ singlet ground states in the strong spin-orbit coupling (SOC) limit and geometric frustration in a face-centered cubic structure formed by the Ir$^{5+}$ ions, we observe this iridate to undergo a novel magnetic transition at temperatures below 1.3 K. We provide compelling experimental and theoretical evidence that the origin of magnetism is in an unusual interplay between strong noncubic crystal fields, local exchange interactions, and “intermediate-strength” SOC. Sr$_2$YIrO$_6$ provides a rare example of the failed dominance of SOC in the iridates.

DOI: 10.1103/PhysRevLett.112.056402

PACS numbers: 71.70.Ej, 71.70.Ch, 75.30.-m

The iridates have become a fertile ground for studies of new physics driven by strong spin-orbit coupling (SOC) that is comparable to the on-site Coulomb ($U$) and crystalline electric field interactions. This unique circumstance creates a delicate balance between interactions that drives complex magnetic and dielectric behaviors and exotic states seldom or never seen in other materials. A profound manifestation of this competition is the novel “$J_{\text{eff}} = 1/2$ Mott state” that was recently observed in the layered iridates with tetravalent Ir$^{4+}(5d^5)$ ions [1–3]. In essence, strong crystal fields split off $5d$ band states with $e_g$ symmetry, whereas the remaining $t_{2g}$ bands form $J_{\text{eff}} = 1/2$ and $J_{\text{eff}} = 3/2$ multiplets via strong SOC. The $J_{\text{eff}} = 3/2$ band is lower in energy and is hence fully filled, leaving the $J_{\text{eff}} = 1/2$ band which is of higher energy half filled. The key, surprising result is the $J_{\text{eff}} = 1/2$ band has a small enough bandwidth that even a modest Coulomb repulsion $U$ among the $5d$-electron states is sufficient to open a Mott gap in these iridates [1,2], which is contrary to expectations based upon the relatively large unperturbed $5d$ bandwidth.

A great deal of recent theoretical and experimental work has appeared in response to early experiments, including predictions of a large array of novel effects in $5d$-electron systems having strong SOC: superconductivity [4], Weyl semimetals with Fermi arcs [5], correlated topological insulators with large gaps, Kitaev spin liquids [6–18], etc. Most of these discussions have focused on the tetravalent iridates in which the Kramers degeneracy of the Ir$^{4+}(5d^5)$ ions results in magnetism. On the other hand, very little attention has been drawn to iridates having pentavalent Ir$^{5+}(5d^4)$ ions, primarily because the strong SOC limit is expected to lead to a nonmagnetic singlet ground state, which can be simply understood as a $J_{\text{eff}} = 0$ state arising from four electrons filling the lower $J_{\text{eff}} = 3/2$ quadruplet [see Fig. 1(c) for a cartoon picture]. Indeed, the $J_{\text{eff}} = 0$ state has been used to explain the absence of magnetic ordering in the pentavalent post perovskite NaIrO$_3$ [19].

An interesting issue that has received some but limited attention is how the $J_{\text{eff}}$ picture is affected by noncubic crystal fields [20]. The presence of such crystal fields has been clearly observed in a number of recent experimental works on materials with the Ir$^{4+}(5d^5)$ electronic configuration such as Sr$_3$IrCuO$_6$ [21], Na$_2$IrO$_3$, and Li$_2$IrO$_3$ [22]. It is generally agreed in these studies that for Ir$^{4+}(5d^5)$, despite the presence of the noncubic crystal field and its importance for the electronic structure, the basic $J_{\text{eff}}$ picture is a good starting point to understand the magnetism in these iridates. In this Letter, we show that, in contrast, for the Ir$^{5+}(5d^4)$ ions of Sr$_2$YIrO$_6$ the strong noncubic crystal field results in a breakdown of the $J_{\text{eff}}$ picture even as a starting point. As illustrated in Fig. 1(c), the conventional strong spin-orbit coupling picture, popular in the description of the Ir$^{4+}(5d^5)$ systems, would result in the ionic state of a single Ir$^{5+}(5d^4)$ being a nonmagnetic singlet, predicting then that Sr$_2$YIrO$_6$ should be a band insulator with no magnetism. Unexpectedly, we find instead from experiment that Sr$_2$YIrO$_6$ hosts well-formed magnetic moments and a magnetic transition below 1.3 K. We shall see that this surprising behavior and other features of the experiment, such as the small amount of entropy lost at the transition, can be understood if we take into account the presence of a substantial noncubic crystal field on the Ir sites. This crystal field arises from distortions of the IrO$_6$ octahedra that are evident from the crystal structure inferred from an x-ray analysis. Finally, we observe unusual metamagnetic behavior at low temperatures whose origin lies in the face-centered cubic (fcc) lattice that the Ir moments form in this ordered double perovskite (see, e.g., [20,23–29]). Quantum
magnetism on the frustrated fcc lattice is yet poorly understood both experimentally and theoretically, and so our work calls for detailed neutron scattering and magnetic x-ray studies of this unusual quantum magnet.

Sr₂YIrO₆ adopts a monoclinic structure essentially derived from the perovskite SrIrO₃ by replacing every other Ir by nonmagnetic Y; the remaining magnetic Ir⁵⁺ ions form a fcc structure with lattice parameters elongated compared to the parent cubic structure, as shown in Fig. 2. Because of the differences in valence state and ionic radius between Y³⁺ and Ir⁵⁺ ions, no significant intersite disorder is expected. The lattice parameters of Sr₂YIrO₆ are given in Table 1 in the Supplemental Material [30]. The IrO₆ octahedra are tilted and rotated, as seen in Fig. 2. A crucial structural detail is that each IrO₆ octahedron is significantly flattened since the bond distance between Ir and apical oxygen Ir-O₃ (= 1.9366 Å) is considerably shorter than the in-plane Ir-O1 and Ir-O2 bond distances (= 1.9798 Å and 1.9723 Å, respectively), as shown in Fig. 2(b). The flattening of the IrO₆ octahedron generates a noncubic crystal field Δ that strongly competes with the spin-orbit interaction λ, as discussed below.

Sr₂YIrO₆ displays paramagnetic behavior at temperatures above 1.5 K, as the magnetic susceptibility χ(T) follows the Curie-Weiss law for 50 < T < 300 K, as shown in Fig. 3(a). Data fits to the Curie-Weiss law over the range 50 < T < 300 K yield an effective moment μ₀eff = 0.91 μ₃B/Ir and a Curie-Weiss temperature θCW = −229 K. The value of μ₀eff is considerably smaller than the value 2.83 μ₃B/Ir expected for a conventional S = 1 5d-electron system. In fact, a reduced value of μ₀eff is commonplace in iridates [20,23,31–34] in part because the strong SOC may suppress the spin moment [35]. A strong antiferromagnetic exchange coupling might be inferred from the large magnitude of θCW (= −229 K); however, the absence of any magnetic ordering at T > 1.5 K indicates the existence of strong quantum fluctuations in Sr₂YIrO₆. A signature for long-range antiferromagnetic order is evident at a very low temperature, Tₙ = 1.3 K, as shown in Fig. 3(b). The two temperature scales evident in the magnetic data yield a strikingly large frustration parameter, |θCW|/Tₙ = 176.2.

The magnetic state undergoes a sharp metamagnetic transition at a critical field Hc below Tₙ, as shown in Fig. 3(c). The isothermal magnetization M(H) initially rises and then exhibits a plateau before a rapid jump at Hc. The metamagnetic transition signals a spin reorientation, but the remarkably low ordered moment (< 0.023 μ₃B/Ir, even at H > Hc) implies that the magnetic state is only partially ordered or unsaturated. Note that a field dependence of M(H) that features a plateau followed by a metamagnetic transition is observed for some geometrically frustrated magnets [36]. The onset of long-range magnetic order is confirmed by an anomaly in the specific heat C(T) observed near Tₙ, as shown in Fig. 4(a). This anomaly is well-defined but weak, and is extremely sensitive to low magnetic fields, as can be seen in Fig. 4(b). The entropy removal S(T) due to the magnetic transition is finite but very small compared to that expected for any possible magnetic ground states consistent with Jₐeff= 1 or 2, or S = 1.
In light of the data presented above, it is clear that neither SOC $\lambda$ nor noncubic crystal field $\Delta$ alone dominates the low-temperature behavior of $\text{Sr}_2\text{YIrO}_6$. Indeed, for $\lambda \gg \Delta$, a prevailing SOC would suppress magnetic order and render a singlet ground state $J_{\text{eff}} = 0$ (Fig. 1c), which is clearly inconsistent with the experimental observation. On the other hand, a $S = 1$ ground state would occur if $\Delta \gg \lambda$ [Fig. 1(b)]. This scenario cannot adequately account for the small amount of entropy lost at the transition. It is therefore compelling to attribute the observed magnetic state to a delicate interplay between the competing $\lambda$ and $\Delta$.

The standard accepted Hamiltonian for electrons in a $t_{2g}$ manifold is given by a sum of one-body terms: $\lambda$, $\Delta$, $U$, and Hund’s rule exchange $J_H$. The physics of this regime can be understood by first diagonalizing the problem with $\Delta$ and $J_H$. In essence, the

$$H_{t_{2g}} = H_{1b} + H_{mb}$$

$$H_{1b} = - \sum_{m,m',s,s'} c_{m's'}^\dagger (\lambda \hat{I}_{mm'} \vec{s}_{ss'} + \Delta (\hat{\mathbf{I}} \cdot \hat{n})_{mm'} \delta_{ss'}) c_{m's'}$$

$$H_{mb} = U \sum_{m,s} c_{ms}^\dagger c_{ms}^\dagger c_{ms} c_{ms}^\dagger + \frac{J_H}{2} \sum_{m,m',s,s'} \left( \sum_{c_{m's'}^\dagger c_{m's'}^\dagger c_{m's} c_{m's}} \right)$$

where $m$ is an index that labels the $yz$, $xz$, $xy$ orbitals. $\hat{I}$ and $\hat{\mathbf{n}}$ are the spin-1 and spin-1/2 Pauli matrices. $\hat{n}$ is a unit vector in the direction of the noncubic distortion or an Ir-O bond. Since the Ir$^{5+}$ ion carries four $5d$ electrons in the $t_{2g}$ orbitals, Eq. (1) becomes a $15 \times 15$ matrix in this space [see Fig. 1(a)]. We ignore $U$ as we compare the quantum states with the same number of particles; thus, we are left with $\lambda$, $\Delta$, and $J_H$. Since the flattening of the IrO$_6$ octahedra (see Fig. 2) renders $\Delta < 0$ and both $\lambda$ and $J_H > 0$, the Ir$^{5+}$ ion always has a nondegenerate ground state independent of the magnitude of the couplings. However, the excitation gap to the lowest doublet becomes substantially suppressed in the regime where all three parameters are comparable, as shown in Fig. 1.

The physics of this regime can be understood by first diagonalizing the problem with $\Delta$ and $J_H$. In essence, the
The interactions yields a local term \( \text{Heisenberg interaction on the fcc lattice.} \) Now, adding hence expect the resulting above, we first consider the exchange interactions in the essence of the magnetism. Following our discussion between Ir moments is possible, a cartoon model can capture the states with four \( \delta < 0 \) splits the \( t_{2g} \) orbitals, leaving the \( a_{1g} \) having lower energy than the \( e_g \) states. Populating the states with four \( 5d \) electrons gives a degeneracy of 6 (three singlets and one triplet); the presence of \( J_H > 0 \) then promotes a robust \( S = 1 \) ground state, as discussed above. Now adding the SOC \( \lambda \) to the interactions splits the triplet, resulting in a singlet ground state and a doublet excited state. This is the near degeneracy, as shown in Fig. 1. If \( \Delta > \lambda \), the splitting, \( \Delta \), can be fairly small.

While a full super-exchange calculation of the interaction between Ir moments is possible, a cartoon model can capture the essence of the magnetism. Following our discussion above, we first consider the exchange interactions in the absence of \( \lambda \), where spin rotation symmetry is preserved. We hence expect the resulting \( S = 1 \) moments to interact with a Heisenberg interaction on the fcc lattice. Now, adding \( \lambda \) to the interactions yields a local term \( \delta \langle S_i^2 \rangle^2 \) on the \( i\)th Ir site. Putting these together, we arrive at the following \( S = 1 \) model \( H_p = J \sum_{ij} \vec{S}_i \cdot \vec{S}_j + \delta \sum_i \langle S_i^2 \rangle^2 \). Such models have been studied extensively on bipartite lattices (although not on the fcc lattice of interest here) both in theory [37] and experiment [38]. The broad feature of such models that is important for our discussion here is that the magnetic order found when \( \delta = 0 \) can persist even when \( \delta > 0 \), albeit with a suppression of the magnetic ordering temperature. As the coupling \( \delta \) is further increased, at a critical value of the ratio \( (\delta/J) \), the magnetic order is completely suppressed at a quantum critical point.

This scenario predicts that a magnetic ordering can occur even though a single ion can be in a nondegenerate "singlet" state. A unique characteristic of such a magnetic order is that the entropy, which is removed at the magnetic transition, is much smaller than the \( R \times \ln (3) = 9.13 \) J/mole K expected for an ordering transition of \( S = 1 \) moments as the isolated ions already lose their entropy when \( T \ll \delta \). This prediction is consistent with the experimental observation that the entropy is only \( \sim 0.01 \) J/mol K, a tiny fraction of \( R \ln(3) \), as shown in Fig. 4(c). Indeed, the ground state is so fragile that even low magnetic fields are strong enough to tip the balance, based upon the data presented herein; this qualitatively explains (a) the strongly depressed, weak magnetic order [Fig. 3(b)], (b) the drastic changes in the specific heat and entropy in weak applied magnetic fields (Fig. 4), and (c) the low-field metamagnetic transition [Fig. 3(c)].

In conclusion, the relative effect of the SOC critically depends upon the strength of noncubic crystal fields, electron hopping and exchange interactions; therefore, it should vary from material to material. While the \( J_{\text{eff}} \) model successfully captures the new physics observed in many iridates, it may not be appropriate to describe new phenomena in other heavy transition metal materials with strong noncubic crystal fields. Our results for \( \text{Sr}_2\text{YIrO}_6 \) not only illustrate a prototypical breakdown of spin-orbit-driven \( J_{\text{eff}} \) states, but also provide a new paradigm for studying novel phenomena emerging from the strong competition between the SOC, local exchange interactions, and crystal fields [39,40].

GC is very thankful to D. I. Khomskii, T. Saha-Dasgupta, G. Jackeli, Y. B. Kim, D. Singh, I. Mazin, S. V. Streltsov, and X. Dai for enlightening discussions. This work was supported by the U.S. National Science Foundation via Grants No. DMR-0856234, No. DMR-1265162, and No. DMR-NSF DMR-1056536 (RKK).

Note Added.—Recently, we became aware of a theoretical Letter just published in Phys. Rev. Lett. by G. Khaliullin [39] who predicts that the magnetic response is dictated by gapped singlet-triplet excitations in Mott insulators with the \( t_{2g} \) electronic configuration such as of \( \text{Ru}^{4+}, \text{Re}^{3+}, \text{Os}^{4+} \), and \( \text{Ir}^{5+} \) ions. Our Letter presents the first experimental evidence of such magnetism.
