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Spin-orbit driven magnetic insulating state with $J_{\text{eff}}=1/2$ character in a 4d oxide

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The role of relativistic spin-orbit coupling (SOC) in creating the diverse behavior of transition metal oxides (TMOs) has largely been considered a perturbation. While this has proven a valid approximation when describing 3d based TMOs the recent intense interest in 5d systems has been driven specifically by the increased effects of SOC [1]. In 5d TMOs SOC competes with increased orbital overlap (altered bandwidth), reduced on-site Coulomb interactions ($U$) and enhanced crystal field splitting. The resulting behavior includes potential realizations of Weyl semi-metals [2, 3], Kitaev physics [4], topological insulators [5] and routes to unconventional superconductivity [6, 7].

One of the most dramatic and well studied manifestations of SOC in 5d systems was first observed in Sr$_2$IrO$_4$ with the creation of a magnetic $J_{\text{eff}}=1/2$ Mott insulating state [8, 9]. This ground state emerges in the Ir$^{4+}$ ion with 5$d^5$ occupancy due to the $t_{2g}$ manifold, in the limit of cubic crystal field splitting, being split by SOC into a filled $J_{\text{eff}}=3/2$ band and a half filled $J_{\text{eff}}=1/2$ band, with even the reduced $U$ in 5d systems being able to drive the opening of the insulating band gap via the Mott mechanism. Subsequent investigations on iridates have led to a growing list of candidate $J_{\text{eff}}=1/2$ magnetic materials [1]. Observations of $J_{\text{eff}}=1/2$ magnetic insulating behavior outside of the iridates is limited, with no known examples in 4d oxide based systems. Of particular interest in SOC enhanced magnetic states is the bond directional anisotropy of the $J_{\text{eff}}=1/2$ pseudo-spins and how alterations from the ideal case, such as tetragonal distortions or pressure-driven increase in bandwidth in Sr$_2$IrO$_4$ that causes mixing with $J_{\text{eff}}=3/2$ bands [10], influences the creation of exotic ground states.

The limit of a pure $J_{\text{eff}}=1/2$ state requires strong SOC and a cubic CEF environment. While 5d iridates fall into the strong SOC regime no current examples have a cubic environment. Nevertheless even with appreciable distortions the state is still realized in 5d systems, albeit with potential mixing in of the $J_{\text{eff}}=3/2$ bands [1]. The apparent robustness of the ground state indicates that the converse would be applicable: $J_{\text{eff}}=1/2$ behavior should be manifested in systems with near ideal octahedra even if the SOC is reduced from those found in 5d systems. However in 4d TMOs where single-ion SOC is of the order 0.15 eV [11, 12], compared to 0.5 eV for iridates, it is unclear whether $J_{\text{eff}}=1/2$-like Mott magnetism is manifested or whether they fall into the class of three Kramers states [12]. Outside oxides, examples of SOC enhanced behavior in 4d systems are limited to theoretical predictions in fluoride based paramagnetic Ir and Rh systems, characteristic behavior in the chloride $\alpha$-RuCl$_3$ and non-magnetic semiconducting behavior in Li$_2$RhO$_3$ [13–16].

Here we show the first 4d based oxide compound to exhibit a magnetically ordered ground state with $J_{\text{eff}}=1/2$ character that offers a unique viewpoint on SOC enhanced behavior in general. The material we focus on, Sr$_4$RhO$_6$, hosts a Rh valence of 4+ (4$d^5$), the same valence as observed in the $J_{\text{eff}}=1/2$ iridates (Ir$^{4+}$, 5$d^5$). Previous investigations of Sr$_4$RhO$_6$ have been extremely limited [17, 18]. Vente et al. observed an anomaly in the susceptibility around 8 K that suggested Sr$_4$RhO$_6$ was the first magnetically ordered Rh$^{4+}$ compound, however no microscopic long range magnetic order has been measured.

Sr$_4$RhO$_6$ forms the same hexagonal space group $R\bar{3}c$ adopted by the $J_{\text{eff}}=1/2$ iridate Ca$_4$IrO$_6$. Although not having an $O_h$ point group Ca$_4$IrO$_6$ was shown to reside close to the local cubic limit required for an unmixed $J_{\text{eff}}=1/2$ state [19]. We directly compare the properties and the role of SOC in these related 4d and 5d materials. Structurally we find the rhodate to reside even closer to the local cubic limit and, since it is isostructural to Ca$_4$IrO$_6$, has disconnected RhO$_6$ octahedra. This makes Sr$_4$RhO$_6$ an appealing candidate to look for $J_{\text{eff}}=1/2$ behavior since the expected narrow bands and strong $10Dq$ splitting will help to overcome the reduced SOC
Polycrystalline samples of Sr$_4$RhO$_6$ were grown following a similar method described in the literature [18]. A stoichiometric mixture of high-purity SrCO$_3$, Rh$_2$O$_3$ were ground, pressed into pellets, and sintered in pure oxygen atmosphere at 900°C, 1000°C, and 1100°C with intermediate grindings. The reaction time for each temperature was 4 days. The sample was finally annealed at 1250°C for 15 days. The extensive heating is required to fully remove the Sr$_2$RhO$_4$, phase. A pellet was measured with a PPMS system to probe the resistivity. Neutron powder diffraction (NPD) measurements were performed on a 0.5g sample on the diffractometer HB-2A at the High Flux Isotope Reactor (HFIR), ORNL. Measurements of the crystal structure were performed with a wavelength of $\lambda=1.54$ Å and measurements of the magnetic structure were performed with $\lambda=2.41$ Å. Temperature measurements at select 20° were performed with the HB-1A fixed incident energy triple-axis spectrometer with $\lambda=2.36$ Å. The branching ratio of the L-edges was measured with x-ray absorption near edge spectroscopy (XANES) at beamline 4-ID-D at the Advanced Photon Source (APS), Argonne National Laboratory. The fluorescence was collected with a detector placed upstream from the sample at 22° from the incoming beam. Measurement were performed on different areas of the powder-on-tape sample to ensure reproducibility. The DFT calculations were performed with the generalized gradient approximation and projector augmented wave (PAW) approach [20] as implemented in the Vienna ab initio simulation package (VASP) [21, 22]. For Rh and O standard potentials were used (Rh and O in the VASP distribution), and for Sr a potential in which semicore s and p states are treated as valence states is used (Srsv). The structural optimization was done using the doubled unit cell with the experimental lattice constants, a 2x2x2 k-point grid and an energy cutoff of 550 eV. Subsequently, the magnetic ground state was examined including the local $U$ for the Rh d states to account for strong correlation effects [23] with $U=2.5$ and $J_H=0.9$ eV, where $J_H$ is the Hund’s coupling [24], as well as the SOC.

The structure of Sr$_4$RhO$_6$ has been previously established by laboratory x-ray measurements [18], however this technique is appreciably less sensitive to oxygen positions compared to neutrons. Given the importance of octahedral distortions in potentially controlling the mixing and subsequent $J_{\text{eff}}=1/2$ character and emergent magnetic and electronic properties we performed NPD on the HB-2A diffractometer. The refined structure based on those in the literature remained stable to 3 K, within the magnetic regime. The sample was found to be of high quality with no Sr$_2$RhO$_4$ that forms in the intermediate growth phase and only $\sim 1.3\%$ non-magnetic SrO impurity phase. The crystal structure obtained from NPD is shown in Fig. 1. The disconnected RhO$_6$ octahedra are evident, with no shared Rh-O bonds between octahedra. Moreover the ideal cubic nature of these octahedra are revealed with identical Rh-O bonds (2.0437(14) Å) and two O-Rh-O bond angles that deviate by a remarkably small amount of less than 0.1° from the ideal 90°. This contrasts with a deviation of $\sim 2°$ in Ca$_4$IrO$_6$ that is considered one of the closest $J_{\text{eff}}=1/2$ with nearly regular octahedra [19]. Therefore structurally Sr$_4$RhO$_6$ appears well suited to host electronic and magnetic behavior that can be controlled by SOC.

Before considering the electronic behavior further we address the significant question of whether Sr$_4$RhO$_6$ orders magnetically and the nature of the ground state.
NPD has proven challenging for $J_{\text{eff}}=1/2$ Ir-based compounds due to the combination of large neutron absorption, small moment size and pronounced magnetic form factor induced reduction in intensity with scattering angle. All of these factors are reduced when moving from Ir to Rh based systems and this allowed a successful NPD investigation with measurements performed through the reported anomaly in susceptibility to probe the microscopic magnetic structure. Comparing results at 15 K and 3 K shows the presence of extra intensity at low temperature in Sr$_4$RhO$_6$. The results of magnetic refinements of the NPD for $\Gamma_1$ are shown in Fig. 2(a). The ordered magnetic moment from NPD measurements is 0.66(5)$\mu_\text{B}$/Rh ion. Following the intensity of one of the magnetic reflections with temperature yields a magnetic ordering temperature of 7.4(5) K, as shown in Fig. 2(b). Contrasting with results on Ca$_4$IrO$_6$ [19] reveals both compounds adopt the same magnetic structure, although with slightly different canting from the c-axis and although the moment sizes are similar it is slightly larger in the rhodate. This would be expected in two compounds with similar magnetic ordering temperatures and underlying behavior, with the principle distinction being reduced itinerancy in the rhodate and subsequent increased local moment.

With the similarities between Sr$_4$RhO$_6$ and the $J_{\text{eff}}=1/2$ iridate Ca$_4$IrO$_6$ in terms of the magnetic structure we now experimentally consider the role of SOC on Sr$_4$RhO$_6$. A useful probe in this regard for the iridates has proven to be x-ray absorption spectroscopy, where...
the branching ratio (BR) of the L-edges provides evidence for the existence of enhanced SOC if the ratio deviates from the statistical value of 2 [26]. The BR is obtained by fitting the white lines from XANES measurements to a resolution broadened step function plus a lorentzian to obtain the integrated intensity, shown as the grey area in Fig. 3. The results reveal BR=3.6(2) for Sr$_4$RhO$_6$. The strong deviation from statistical indicates an appreciable role for SOC in the ground state. Indeed the values correspond closely to the BR for iridates that show strongly SOC enhanced behavior [26]. Similar XANES measurements on α-RuCl$_3$ revealed a BR of 3 that was presented as evidence for substantial SOC [14].

To further investigate the role of SOC we performed DFT calculations on Sr$_4$RhO$_6$ including the $Γ_1$ magnetic ground state. The results (Fig. 4(a)) show a $t_{2g}$ manifold split into $J_{\text{eff}}=1/2$ and $J_{\text{eff}}=3/2$ dominated bands. The $J_{\text{eff}}=1/2$ states are higher in energy then $J_{\text{eff}}=3/2$ with the near-Fermi-level states given predominantly by the $J_{\text{eff}}=1/2$ states, and $J_{\text{eff}}=3/2$ fully filled. Therefore, while the splitting between $J_{\text{eff}}=1/2$ and $J_{\text{eff}}=3/2$ is not complete, as can be seen in the DOS, spectroscopically a pure $J_{\text{eff}}=1/2$ is realized because the scattering process is from the excited to the unoccupied states. Experimentally we probed the insulating nature with resistance measurements on a pressed pellet, see Fig. 4(b). These results are consistent with an insulating material and moreover the powder is dull, characteristic of an insulator. From DFT calculations an insulating gap of around 0.1 eV is observed. This insulating gap is reduced from that found in Ca$_4$IrO$_6$ from DFT of 0.6 eV. Moreover we find that unlike Ca$_4$IrO$_6$ the $J_{\text{eff}}=1/2$ and $J_{\text{eff}}=3/2$ bands are not fully separated. This mirrors the situation from DFT calculations in the paramagnetic fluoride based Ir and Rh systems predicted to both be $J_{\text{eff}}=1/2$ Mott insulators [13]. The mixed $J_{\text{eff}}$ bands indicates that despite the essential ideal octahedra in Sr$_4$RhO$_6$ the intrinsic SOC is not large enough to fully split the $t_{2g}$ manifold. This situation is analogous to several other candidate $J_{\text{eff}}=1/2$ materials where the opposite occurs: the $J_{\text{eff}}=1/2$ and $J_{\text{eff}}=3/2$ bands are mixed despite the large SOC due to appreciable non-cubic distortions.

Collectively our experimental and theoretical results reveal the behavior in Sr$_4$RhO$_6$ is strongly influenced by SOC. Qualitatively the results are very similar to the Rh based fluorides, however it is important to consider the differing effects of oxygen compared to fluorine. Specifically in oxides the hybridization is generally much stronger than fluorides. Consequently the band width of $J_{\text{eff}}=1/2$ dominated bands and $J_{\text{eff}}=3/2$ dominated bands are narrower in fluorides, of the order 0.1 eV compared to 0.5 eV in Sr$_4$RhO$_6$. Therefore while the degree of separation is stronger in the fluorides both show $J_{\text{eff}}=1/2$ character, although neither show fully separated $J_{\text{eff}}=1/2$ and $J_{\text{eff}}=3/2$ dominated bands. This leads to the question of whether an alternative approach is more appropriate in which both these fluoride and oxides system are best described by three Kramers states as considered for other rhodates [12]. However, as suggested in Ref. 12 the BR from XANES should distinguish between these cases and indeed this does so in Sr$_4$RhO$_6$.

In conclusion, we have observed a rare occurrence of long range magnetic order in a rhodate compound. Moreover Sr$_4$RhO$_6$ is found to be an insulator leading to a consideration in terms of SOC enhanced behavior. Experimentally the magnetic ordering and insulating behavior are all analogous to the isostuctural $J_{\text{eff}}=1/2$ iridate Ca$_4$IrO$_6$. DFT calculations reveal $J_{\text{eff}}$ character with mixed $J_{\text{eff}}=1/2$ and $J_{\text{eff}}=3/2$ dominated bands. Despite the mixing due to the reduced SOC in going from Ir to Rh the physical properties are strongly influenced by SOC. Therefore while this is the first 4d TMO to be shown to have $J_{\text{eff}}$ character further investigations on oxides with similar spatially disconnected octahedra offer routes to uncovering analogous exotic properties as found in the iridates.

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