

Magnetic Structures of SrIrO₃/SrTiO₃ Superlattices Studied by Resonant X-Ray Diffraction

5d Ir oxides have an interplay of spin-orbit coupling and electron correlations. We investigated the magnetic structure of a SrIrO₃/SrTiO₃ superlattice by resonant X-ray diffraction. We observed the (0.5, 0.5, 5) peak, which indicates in-plane antiferromagnetic ordering and interlayer ferromagnetic coupling. Together with the weak ferromagnetic moment in the superlattice observed only for magnetic field parallel to the IrO₂ plane, we concluded that in-plane canted antiferromagnetism is realized in this superlattice. These results show that we can design and realize novel electronic phases in the unit of the SrIrO₃ monolayer.

The novel interplay of spin-orbit coupling (SOC) and electron correlations in complex Ir oxides (iridates) has recently emerged as a new paradigm for correlated electron physics. The transition from semimetallic SrIrO₃ to magnetic insulator Sr₂IrO₄ is reported in Ruddlesden-Popper series Sr_{n+1}Ir_nO_{3n+1} ($n = 1, 2, \text{ and } \infty$), suggesting a dimensionality controlled bandwidth via n [1]. Matsuno *et al.* reported transport and magnetic behaviors of synthesized artificial superlattice [(SrIrO₃) _{m} , SrTiO₃] with $m = 1, 2, 3, 4, \text{ and } \infty$ [2]. We tracked the evolution of electronic ground states by varying m to understand the close links among semimetal-insulator transition, magnetism, and underlying lattice structures. **Figure 1a** shows the schematics of the superlattices [(SrIrO₃) _{m} , SrTiO₃] ($m = 1, 2, \text{ and } \infty$). In this work, we investigated the magnetic structure of the SrIrO₃/SrTiO₃ ($m = 1$) superlattice by resonant X-ray diffraction to clarify the exact magnetic structure [2].

The SrIrO₃/SrTiO₃ superlattice sample was fabricated on SrTiO₃(001) substrate by a pulsed laser deposition (PLD) technique. The details of the sample fabrication were described elsewhere [2]. Resonant X-ray diffraction measurements were performed at beamline 3A at Photon Factory, KEK. Photon polarization of the $\sigma\text{-}\pi'$ channel was selectively measured by using a Mo(400) analyzer crystal.

Figure 1b shows the temperature dependence of the magnetization measured at 0.1 T for the SrIrO₃/SrTiO₃ superlattice. The weak ferromagnetic moment in the superlattice was indeed observed only for magnetic field parallel to the IrO₂ plane. This is consistent with the idea that weak moments originate from the Dzyaloshinskii-Moriya (DM) interaction associated with the in-plane rotation of IrO₆ octahedra. **Figure 1c** shows the temperature dependence of the magnetic X-ray diffraction intensity for the (0.5, 0.5, 5) peak measured at Ir L_3 edge with $\sigma\text{-}\pi'$ polarization to extract the magnetic contribution [3]. Here, the unit cell dimensions are $a \times a \times 2a$ ($a \sim 0.39$ nm). This peak shows the in-plane antiferromagnetic ordering. From these results, we conclude

that an in-plane canted antiferromagnetism is realized in this superlattice, similar to the single layer perovskite Sr₂IrO₄.

Since magnetic diffraction with the integer c -axis index (0.5, 0.5, 5) was observed, the interlayer coupling of Ir local moments is ferromagnetic, in marked contrast to the bulk Sr₂IrO₄. This is naturally expected because the interlayer coupling of Ir moments through the hybridization with the inserted SrTiO₃ layer should give rise to a ferromagnetic Ir-Ir coupling regardless of whether the Ir-Ti coupling is ferromagnetic or antiferromagnetic. **Figure 1d** shows the obtained magnetic and lattice ordering pattern of this superlattice. Here, the IrO₆ octahedra in the two adjacent IrO₂ layers rotate in the same direction, which can account for ferromagnetic coupling of both canted moments and Ir moments.

The canted moment observed for the SrIrO₃/SrTiO₃ superlattice was 0.02 μ_B /Ir mol. This is $\sim 1/4$ of the 0.075 μ_B /Ir mol value in the bulk Sr₂IrO₄ [4], which implies the reduction of the local Ir moment and/or the reduction of DM interaction. The in-plane lattice constant $a \sim 0.3905$ nm for the superlattice sample is larger than the 0.3890 nm value for Sr₂IrO₄ [4] and therefore a smaller rotation of the IrO₆ octahedra is expected for the superlattice. We estimated a rotation angle of approximately 8° in the superlattice. The Ir-O-Ir bond closer to 180° in the superlattice should decrease the strength of the DM interaction and increase the bandwidth. The latter makes the system more itinerant and reduces the magnitude of the magnetic moments. We therefore conclude that the reduced distortion of the lattice is responsible for the reduced canted moments in the superlattice.

In summary, we performed a resonant X-ray diffraction study of the SrIrO₃/SrTiO₃ superlattice to determine the magnetic structure. We observed in-plane canted antiferromagnetism similar to the bulk Sr₂IrO₄, and ferromagnetic interlayer coupling in contrast to Sr₂IrO₄. These results show that we can design and realize novel electronic phases in the unit of the SrIrO₃ monolayer.

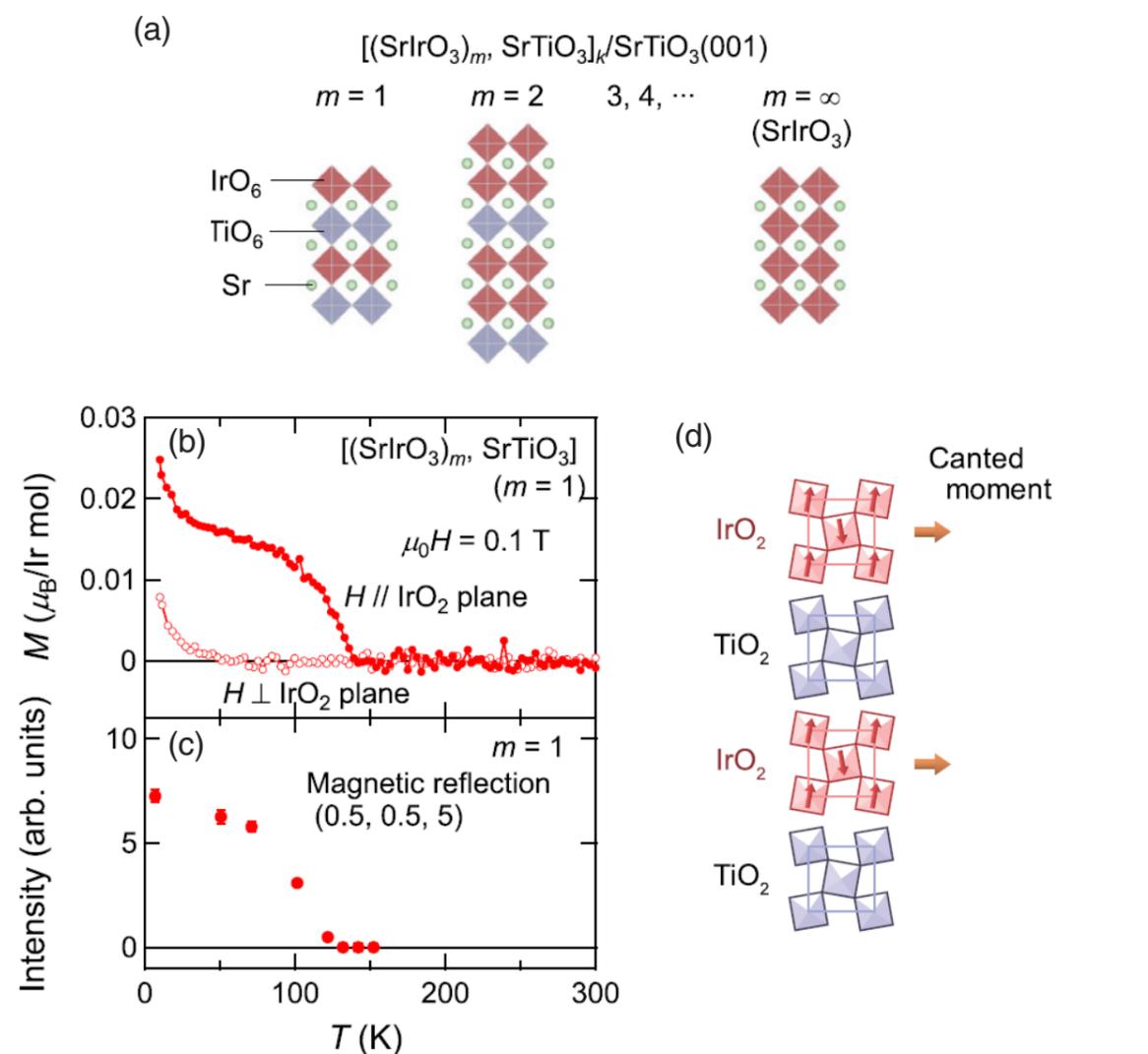


Figure 1: (a) Schematics of the superlattices [(SrIrO₃) _{m} , SrTiO₃] ($m = 1, 2, \text{ and } \infty$). (b) Temperature dependence of the magnetization measured at 0.1 T for the SrIrO₃/SrTiO₃ superlattice. (c) Temperature dependence of the magnetic X-ray diffraction intensity for the (0.5, 0.5, 5) peak. (d) Magnetic structures of the SrIrO₃/SrTiO₃ superlattice.

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