Off-stoichiometry effects on the structural and transport properties of epitaxial $SrRu_xO_3$ (x = 0.7 and 1) thin films

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The itinerant 4*d* ferromagnetic perovskite SrRuO₃ has been widely used in oxide electronics and spintronics as an epitaxial conducting layer [1]. Recently, renewed interest on SrRuO₃ emerged after the discovery of Weyl fermions in SrRuO₃ [2-4]. The quantum transport of the Weyl fermions can be detected only for samples having a high residual resistivity ratio (RRR) [3.4]. In general, it is possible that the Ru stoichiometry may play a role in realizing the magnetic Weyl states [1]. However, there is no unified understanding on the difference in magnetic and transport properties between SrRuO₃ films grown under stoichiometric and off-stoichiometric conditions.

In this presentation, we report the thickness (*t*)-dependent magnetotransport properties of epitaxial SrRu_xO₃ (x = 0.7 and 1) films grown on SrTiO₃ substrates by molecular beam epitaxy [5]. For the films with x = 0.7, the Ru deficiency was intentionally introduced by varying the supplied Ru/Sr flux ratio. Notably, the crystalline quality for such off-stoichiometry films is comparable for x = 0.7 and 1, and structural characterizations using diffraction or microscopy techniques can hardly distinguish any differences [5]. The residual resistivity ρ_{Res} of the SrRu_{0.7}O₃ film is much higher than that of SrRuO₃ at each *t* [Fig. 1(a)], indicating that the Ru vacancies serve as carrier scattering mechanism due to interface-driven disorders exists. Despite the large off-stoichiometry, SrRu_{0.7}O₃ ($t \ge 10$ nm) shows a Curie temperature (T_{C}) of ~ 140 K, which is only ~10 K lower than that in SrRuO₃ [Fig.

1(b)]. For t < 10 nm, $T_{\rm C}$ abruptly decreases with decreasing t. We conclude that the interface-driven disorder has a detrimental influence on the magnetic exchange interaction. In addition, the Weyl transport is not observed in the off-stoichiometric SrRu_{0.7}O₃ films. Therefore, proper stoichiometry control is vitally important to utilize the full potential of SrRuO₃ as a magnetic Weyl semimetal [5].



Figure 1 (a) Thickness *t* dependence of the residual resistivity ρ_{Res} of the stoichiometric (SrRuO₃) and Ru-deficient SrRu_{0.7}O₃ films. (b) Thickness *t* dependence of the *T*_C.

References [1] G. Koster *et al.*, Rev. Mod. Phys. **84**, 253 (2012). [2] Y. K. Wakabayashi, *et al.*, APL Mater. **7**, 101114 (2019). [3] K. Takiguchi, Y. K. Wakabayashi, *et al.*, Nat. Commun. **11**, 4969 (2020). [4] S. Kaneta-Takada, Y. K. Wakabayashi, *et al.*, Appl. Phys. Lett. **118**, 092408 (2021). [5] Y. K. Wakabayashi, *et al.*, AIP Advances **11**, 035226 (2021).