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Size Effect on the Magnetic Phase in Sr₄Ru₃O₁₀

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Liu, Yan; Yang, Jiyong; Wang, Weike; Du, Haifeng; Ning, Wei; Ling, Langsheng; Tong, Wei; Qu, Zhe; Yang, Zhaorong; Tian, Minling; Cao, Gang; and Zhang, Yuheng, "Size Effect on the Magnetic Phase in $Sr_4Ru_3O_{10}$ " (2016). *Center for Advanced Materials Faculty Publications*. 4.

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Notes/Citation Information

Published in New Journal of Physics, v. 18, article 053019, p. 1-6.

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Digital Object Identifier (DOI)

https://doi.org/10.1088/1367-2630/18/5/053019

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Size effect on the magnetic phase in $Sr_4Ru_3O_{10}$

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Keywords: Sr₄Ru₃O₁₀, metamagnetic transition, size effect, Hall effect

Abstract

High quality $Sr_4Ru_3O_{10}$ nanoflakes are obtained by the scotch tape-based micro-mechanical exfoliation method. The metamagnetic transition temperature T_m flake is found to decrease in line with the decrease of thickness, while the ferromagnetic (FM) phase, the ordinary, and anomalous Hall effects (OHE and AHE) are independent on the thickness of the flake. Analysis of the data demonstrates that the AHE reflects the FM nature of $Sr_4Ru_3O_{10}$, and the decrease of thickness favors the Ru moments aligned in the *ab*-plane, which induces a decrease of the metamagnetic transition temperature compared with the bulk.

1. Introduction

Ruthenium oxide perovskites $Sr_{n+1}Ru_nO_{3n+1}$ ($n = 1, 2, 3, \infty$) are strongly correlated materials involving complex interactions between the charge, spin, orbit and lattice degrees of freedom. Their ground states present a rich of exotic physical properties, such as the unconventional spin-triplet superconductivity in Sr₂RuO₄ (n = 1) [1], the quantum criticality and nematicity in Sr₃Ru₂O₇ (n = 2) [2–4], and the spontaneous itinerant ferromagnetism in the infinite layer of SrRuO₃ ($n = \infty$) [5, 6]. Sr₄Ru₃O₁₀ is the n = 3 member of the $Sr_{n+1}Ru_nO_{3n+1}$ family. It belongs to quasi-two-dimensional metal with an orthorhombic unit cell, which is composed of triple layers of corner-shared RuO₆ octahedra separated by double rock-salt Sr-O layer [7]. This member has attracted considerable attention in recent years due to its unique magnetic properties [7-17]. By applying a magnetic field H(0.01T) along the c-axis, the temperature dependent magnetization $M_c(T)$ exhibits a FM transition at a Curie temperature $T_c \sim 105$ K, followed by another sharp transition at temperature T_m^{bulk} of about \sim 50 K with a large irreversibility. In contrast, by applying a field (H = 0.01 T) within the *ab*-plane, the $M_{ab}(T)$ displays a weak cusp at T_c and a pronounced peak at T_m^{bulk} with a much smaller irreversibility, and below about 20 K, the M_{ab} is almost unmeasurable [7, 8]. Interestingly, if the in-plane field reaches a critical field H_{co} a rapid increase of Mab in a narrow range of magnetic field is observed on the M-H curves at temperatures below T_m^{bulk} , and the H_c increases with the decrease of temperature $(H_c^{2K} \sim 2 \text{ T})$ [8]. This transition is considered to be the metamagnetic transition.

The metamagnetic transition observed in $Sr_4Ru_3O_{10}$ below T_m^{bulk} contains rich physics associated with the lattice, spin, orbit, and electronic inhomogeneity, which has not been well understood. Firstly, this transition is accompanied by strong spin-lattice coupling [9, 14, 18]. The crystal structure undergoes a significant change when the applied in-plane field crosses over the critical field H_c [9, 14]. Secondly, the occurrence of the transition is possibly through a phase separation process with magnetic domain formation, for the in-plane magnetoresistivity near H_c exhibits large hysteresis or multiple ultra-sharp steps at extreme low temperatures (<1 K) [10, 11, 17]. Thirdly, the angle-resolved magnetization and magnetoresistivity suggest the metamagnetic transition is orbit-dependent, where the Ru $4d_{xz,yz}$ orbit is responsible for the metamagnetic transition while the $4d_{xy}$ orbit is ferromagnetic in the ground state [12, 13]. The understanding of the metamagnetic transition in

PAPER

ACCEPTED FOR PUBLICATION 26 April 2016 PUBLISHED 12 May 2016

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Figure 1. (a) Magnetization as a function of temperature measured along the *c*-axis at 0.01 T of the Sr₄Ru₃O₁₀ single crystal under zero-/or field cooling condition. (b) SEM image of a Sr₄Ru₃O₁₀ flake (d = 260 nm) with patterned electrodes. (c) and (d) are, respectively, the temperature dependence of the longitudinal resistivity ρ_{xx} of the Sr₄Ru₃O₁₀ flakes with different thickness *d*; the insets are, respectively, the blow-up of the ρ_{xx} -*T* curves in low-*T* range measured under in-plane fields.

 $Sr_4Ru_3O_{10}$ is challenged by recent neutron study, where the Ru moments at zero field and below T_c are found to be FM-aligned along the *c*-axis only, there is no any signature of either long-range antiferromagnetic (AFM) or FM order in the *ab*-plane [14]. However, the neutron data cannot exclude a possibility that the metamagnetism is a field-induced short-range AFM order to FM transition. To date, the magnetic nature of $Sr_4Ru_3O_{10}$ below T_m^{bulk} is still unclear and remains elusive.

In this work, we present the electrical transport measurements of two mechanically exfoliated Sr₄Ru₃O₁₀ nanoflakes with thickness of 31 nm and 260 nm. It finds that the metamagnetic transition temperature, T_m flake, in the flake is much smaller than the bulk, T_m bulk ~ 50 K, and decreases with decreasing thickness, but its saturation field H_s along the *c*-axis increases with the decrease of thickness, indicating the Ru moment in the thinner sample is more difficult to be aligned along the *c*-direction. However, the FM transition, the ordinary and anomalous Hall effects are independent on the thickness, where the dominant carriers derived from the ordinary Hall coefficient R_0 are always hole-type and the anomalous Hall conductivity σ_{xy} ^A follows the typical scaling law σ_{xy} ^A $\propto \sigma_{xx}$ φ (σ_{xx} : longitudinal conductivity) with different scaling exponent, $\varphi \sim 1$, 0 and 1.6, as the increase of the unit cell or the electronic structures, but can be understood by the shape anisotropy induced rearrangement of the Ru moments due to the size effect.

2. Experimental

 $Sr_4Ru_3O_{10}$ single crystal is grown by the flux technique [8]. The temperature dependent magnetization of the bulk crystal along the *c*-axis measured under 0.01 T is shown in figure 1(a), which is well consistent with those reported previously [7, 8]. The characteristic temperatures $T_m^{bulk} \sim 50$ K and $T_c \sim 105$ K of the single crystal are indicated by the arrows. The $Sr_4Ru_3O_{10}$ flakes are obtained by the scotch tape-based micro-mechanical exfoliation from the bulk single crystal, and then transferred to a silicon substrate covered with 300 nm-thick silicon dioxide on the top of the surface. The thickness, *d*, of the flake is determined by atomic force microscopy. Conventional six terminal electrical contacts are made using electron-beam lithography (EBL) technique. To



Figure 2. The magnetoresistivity MR of Sr₄Ru₃O₁₀ flakes as a function of temperature *T* under various in-plane magnetic fields. (a) d = 31 nm; (b) d = 260 nm.

ensure ohmic contacts, we etch the e-beam patterned contact areas for a few seconds by *in situ* Ar-plasma to remove the possible residual PMMA resist firstly, and then deposit 5 nm Ti and 150 nm Au working as the electrical leads by thermal evaporation. With this process, the successful rate for ohmic contacts is higher than 90%, and the resistance ratio between the contact resistance and the sample is usually less than 15%. The scanning electron microscope (SEM) image of a device with patterned electrode is shown in figure 1(b). The longitudinal resistivity (ρ_{xx}) and the Hall resistivity (ρ_{xy}) are measured as functions of temperature (T) and external magnetic field (H) by a physical property measurement system (PPMS, Quantum Design). The ρ_{xx} is measured by the standard four-probe configuration, where the contact resistance is believed to be negligible. The Hall resistance ρ_{xy} is determined from $\rho_{xy} = [\rho_{xy}(H) - \rho_{xy}(-H)]/2$ in order to subtract the longitudinal component of ρ_{xx} arising from the small misalignment of the transverse contacts.

3. Results and discussion

Figures 1(c) and (d) show ρ_{xx} versus T curves of two flakes (d = 31 and 260 nm) measured at zero magnetic field. For each flake, the ρ_{xx} decreases monotonically as T decreases. The residual resistance ratio, $RRR = \rho_{xx}^{300 \text{ K}} / \rho_{xx}^{2 \text{ K}}$, of each flake reaches about 60 (data not shown), indicating the crystals are high quality, where the residual resistivity of both flakes at 2 K is about 3.77 $\mu\Omega$.cm (d = 31 nm) or 4.15 $\mu\Omega$.cm (d = 260 nm). Two anomalies on the resistivity can be clearly identified from the magnetoresistivity (MR) measured under various in-plane magnetic fields H as shown in figure 2, where MR is defined as $[\rho_{xx}(H)-\rho_{xx}(0)]/\rho_{xx}(0)$. The data is obtained by the zero field cooling from above 160 K to exclude the influence of the field history. The 'valley' observed on the MR-T curves is as expected due to the itinerant FM nature of $Sr_4Ru_3O_{10}[8]$, which is caused by the suppression of carrier scattering from spin fluctuations under in-plane magnetic field [19]. The T_c is determined to be ~106 K and 101 K for the 31 nm- and 260 nm-thick flakes, respectively, from the negative maximum of the 'valley' [19], which is almost independent on the thickness and consistent with the bulk. With decreasing T below T_c , the Sr₄Ru₃O₁₀ undergoes another transition called the metamagnetic transition at about T_m flake ~ 25 K and 37 K for the 31 nm- and 260 nm-thick flakes, below T_m flake the MR shows a positive behavior, while above which the MR is always negative due to the FM nature of Sr₄Ru₃O₁₀. This feature is consistent with that reported in the bulk [11], except for the T_m flake is almost 20 K lower than the T_m bulk ~ 50 K. It finds that the metamagnetic transition temperature decreases with decreasing the thickness of the flake, while the T_c remains unchanged. Previously, both the electronic and lattice structures have been suggested to be the driving force for the formation of the metamagnetic phase [7, 12–15]. A fact that our sample preparation process does not make any variations of the electronic or lattice structures provides an indication that the metamagnetic transition in Sr₄Ru₃O₁₀ should have an origin of internal magnetic orders, which maybe changed as the decrease of thickness. To understand the mechanism in detail, we performed the Hall effect study on the flakes.

Figure 3 shows the *H*-dependence of ρ_{xy} at various temperatures, where *H* is applied perpendicular to the *ab*plane of the flake. As a FM nature of Sr₄Ru₃O₁₀ below T_c , both the OHE and AHE make contributions to the total Hall resistivity [20], characterized by the 'knee' profile as shown in figure 3. At temperatures far above T_c , the







OHE resistivity $\rho_{xy}{}^o$ governs the total ρ_{xy} with a linear H dependence of $\rho_{xy}{}^o = R_0 H$ (e.g. see the dashed line in figure 3(d)), which is due to the orbital effect of H on the carriers, here R_0 is the ordinary Hall coefficient. With decreasing T, the AHE gradually dominates the ρ_{xy} , and the AHE resistivity $\rho_{xy}{}^A$ reaches a maximum at about 80 K. Then $\rho_{xy}{}^A$ decreases rapidly with the further decrease of T and becomes extremely small at 10 K but can still be recognized (e.g. see the dashed line in figure 3(c)). In spite of the FM-induced hysteresis behavior in the low field range, the trace of the observed ρ_{xy} at a fixed temperature can be well described by the equation of $\rho_{xy} = R_0 H + \rho_{xy}{}^A$, which contains both the OHE and AHE components. The ordinary Hall coefficient R_0 can be extracted from the high field slope, *i.e.* $d\rho_{xy}/dH$ for H = 5 T, of the Hall isotherm, and the $\rho_{xy}{}^A$ below T_c can be simultaneously obtained by extrapolating the linear term to the zero field.

The ordinary Hall coefficient R_0 as a function of temperature is shown in figure 4(a). R_0 is positive in the whole temperature range, indicating the dominant carriers in $Sr_4Ru_3O_{10}$ are holes. The R_0 -T curves for both flakes present almost a same 'S-shaped'-like behavior. Specifically, the $R_0(T)$ shows a maximum near $T_1 \sim 22$ K and a minimum near $T_2 \sim 60$ K below T_c , and these kinks show no direct correlation with the magnetic structures, such as the metamagnetic transition near T_m flake and the FM transition at T_c . Generally, the temperature dependent $R_0(T)$ in quasi-two-dimensional metal is related to the changes of the wave vector dependent electron mean free path or the reconstruction of the Fermi surface [21, 22]. The almost identical $R_0(T)$ in both flakes implies that the electronic structures are almost independent on the thickness of the flakes as expected. We noticed that both $R_0(T)$ display a similar T-dependent feature as the variations of the lattice



parameter *c* determined by the neutron diffraction study [14], indicating that the variations of R_0 in Sr₄Ru₃O₁₀ might be closely connected to this structural change.

Now we turn to the AHE component and use the universal scaling relation [20], $\sigma_{xy}^{A} \propto \sigma_{xx}^{\varphi}$, to uncover the physical implication below T_c (~100 K) in the flakes, where σ_{xy}^A and σ_{xx} are, respectively, the anomalous Hall and longitudinal conductivity, and φ is the scaling exponent. Here, $\sigma_{xy}{}^A$ is estimated by $\sigma_{xy}{}^{A} = \rho_{xy}{}^{A} / [\rho_{xx}{}^{2} + (\rho_{xy}{}^{A})^{2}] \approx \rho_{xy}{}^{A} / \rho_{xx}{}^{2}$ since $\rho_{xy}{}^{A}$ is at least two orders smaller than ρ_{xx} in our case. As shown in figure 4(b), the σ_{xy} ^A is also independent on the thickness of the flake, and can be categorized into three regions labeled as I, II and III with an exponent of $\varphi \sim 1.08$ in T < 23 K (or $\sigma_{xx} > 7.5 \times 10^4$ S cm⁻¹), $\varphi \sim 0$ with $\sigma_{xy}{}^{A} = 235 \text{ S cm}^{-1} \text{ in } 23 \text{ K} < T < 55 \text{ K} \text{ (or } 2.2 \times 10^{4} \text{ S cm}^{-1} < \sigma_{xx} < 7.5 \times 10^{4} \text{ S cm}^{-1} \text{) and } \varphi \sim 1.61 \text{ or } 1.72 \text{ in } T > 55 \text{ K} \text{ (or } \sigma_{xx} < 2.2 \times 10^{4} \text{ S cm}^{-1} \text{), respectively. It is well-known that both the extrinsic and intrinsic$ mechanisms are proposed for the AHE [20], in which the $\sigma_{xy}^{A} \propto \sigma_{xx}$ ($\varphi \sim 1$) in region I indicates the extrinsic skew-scattering mechanism is dominant based on the asymmetric impurity scattering caused by the spin-orbit interaction (SOI), while $\varphi \sim 0$ with a large value of σ_{xy} ^A in region II suggests the intrinsic Berry-phase contribution governs the AHE [23]. Recently, Onoda et al (2008) have developed a unified theory of the AHE in ferromagnets and proposed three scaling regions as a function of the carrier lifetime τ related Born scattering amplitude \hbar/τ (\hbar : Planck constant). For $\hbar/\tau < \sim E_{SO}(E_{SO}: \text{spin-orbit interaction energy}), <math>\varphi = 1$, the extrinsic skew-scattering arising from the vertex correction yields a dominant contribution; For $\sim E_{SO} < \hbar/\tau < E_{FS} \varphi = 0$, σ_{xy} ^A becomes insensitive to the scattering strength because of the intrinsic dissipationless topological Berryphase contribution. Further increasing \hbar/τ , $\varphi = 1.6$ [23, 24]. Usually, it is difficult to observe all the proposed regions in one ferromagnetic sample, possible due to a small variation of \hbar/τ [20, 25]. The scaling behavior in a $Sr_4Ru_3O_{10}$ flake can present all the three scaling regions indicates there is a large change of \hbar/τ in this material. The reason for such change is possibly attributed to the large variation of the structures in $Sr_4Ru_3O_{10}$ as well [14], for the two crossovers, one at about 23 K and the other at about 55 K, among the three scaling regions match well with the temperatures T_1 and T_2 shown in figure 4(a). In other words, the AHE reflects the FM nature of the flake below T_{c} and which is independent on the thickness.

Figure 5 shows the saturation field H_s along the *c*-axis of the two flakes determined from the ρ_{xy} -H curves (see figure 3), where the inset representatively displays an enlarged ρ_{xy} -H curve of the 260 nm-thick flake measured at 40 K, and the H_s is 0.65 T indicated by the arrow. The H_s increases as the decrease of thickness at a fixed temperature, indicating that the magnetic moments in the thinner flake are more difficult to be polarized along the *c*-axis. In other words, the shape anisotropy caused by the decrease of thickness favors the magnetic moments aligned in the *ab*-plane (possibly arranged to be a weak FM order). This assignment is consistent with the observation of the metamagnetic transition T_m flake decreases with the decrease of thickness (see figure 2), where the in-plane magnetic order stabilized by the reduced thickness will suppress the metamagnetic phase.

4. Conclusions

In summary, we have investigated the transport property of two mechanically exfoliated $Sr_4Ru_3O_{10}$ nanoflakes. The result shows that the change of thickness has little effect on the FM phase, the OHE and AHE, but can modulate the metamagnetic transition temperature T_m flake significantly, which is about 25 K for the 31 nmthick flake and 37 K for the 260 nm-thick flake, respectively. The identical Hall effect in the two flakes suggests the decrease of T_m flake cannot be attributed to the changes of the unit cell or the electronic structure. However, we have found that the saturation field along the *c*-axis in the two flakes increases as the decrease of thickness, indicating the Ru moment is aligned closer to the *ab*-plane in the thinner flake due to the size effect, and the metamagnetic transition in $Sr_4Ru_3O_{10}$ is thus caused by a rearrangement of the Ru moments.

Acknowledgments

This work was supported by the National Natural Science Foundation of China (Grant No. 11174294, 11174291, 11374302, 11304319, U1332209, U1332139, U1432251 and U1532153), and the program of Users with Excellence, the Hefei Science Center of CAS and the CAS/SAFEA international partnership program for creative research teams of China.

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