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2007 J. Phys.: Condens. Matter 19 145222

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## Anisotropy dependence of anomalous Hall effect in canonical spin glass alloys

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Received 29 August 2006

Published 23 March 2007

Online at [stacks.iop.org/JPhysCM/19/145222](http://stacks.iop.org/JPhysCM/19/145222)

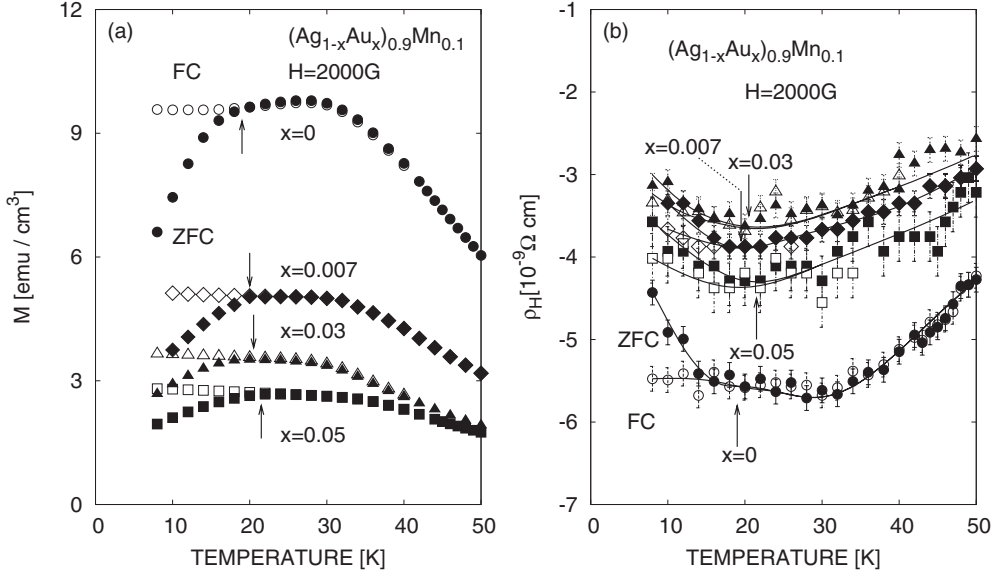
### Abstract

The influence of the Dzyaloshinsky–Moriya (DM) anisotropy on the extraordinary Hall coefficient  $R_s \equiv \rho_{ex}/M$ , where  $\rho_{ex}$  is the extraordinary Hall resistivity and  $M$  is the magnetization, is investigated in canonical spin-glass (SG) alloys. The strength of the DM anisotropy in the alloys is changed systematically by doping with a third impurity that is non-magnetic. The Hall resistivity  $\rho_H$ , the magnetization  $M$  and the resistivity  $\rho$  were measured in the series of  $(\text{Ag}_{1-x}\text{Au}_x)_{0.9}\text{Mn}_{0.1}$  alloys with  $x = 0, 0.007, 0.03, \text{ and } 0.05$ . The difference  $\Delta R_s$  between the values of zero-field-cooled and field-cooled  $R_s$ , below the SG transition temperature, clearly increased with the amount of Au impurities. The dependence of the chirality contribution to  $R_s$  on the DM anisotropy is discussed in relation to the theoretical work for the chirality-driven anomalous Hall effect in the weak coupling regime.

For many decades, the accepted parameterization of the Hall resistivity  $\rho_H$  in magnetic materials has been in terms of the canonical expression

$$\rho_H = \rho_{\text{ord}} + \rho_{\text{ex}} = R_0 H + 4\pi R_s M, \quad (1)$$

where  $\rho_{\text{ord}}$  and  $\rho_{\text{ex}}$  are the ordinary and the extraordinary Hall resistivity respectively,  $R_0$  and  $R_s$  are the ordinary and the extraordinary Hall coefficient respectively,  $H$  is the magnetic field and  $M$  is the magnetization. Recently, some features [1, 2] of  $R_s$  have been reported in canonical spin-glass (SG) alloys which are not understood by the conventional theory [3]. This behaviour of  $R_s$  indicates the existence of the chirality-driven extraordinary Hall effect term as predicted by the theories of the chirality mechanism of the Hall effect [4, 5]. These theories also predict that the Dzyaloshinsky–Moriya (DM) anisotropy plays an important role in the appearance of the chirality-driven extraordinary Hall effect term. The strength of the DM anisotropy of the canonical SG alloys is changed systematically by doping with a third impurity that is non-magnetic. The main purpose of the present article is to investigate how the DM anisotropy



**Figure 1.** (a) Temperature dependence of  $M$  for  $(\text{Ag}_{1-x}\text{Au}_x)_{0.9}\text{Mn}_{0.1}$  in a field of 2000 G. (b) Temperature dependence of  $\rho_H$  for  $(\text{Ag}_{1-x}\text{Au}_x)_{0.9}\text{Mn}_{0.1}$  in a field of 2000 G. The arrows mark  $T_g$  (2000 G).

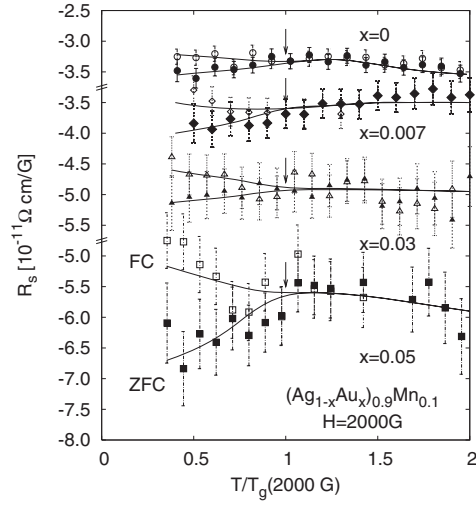
**Table 1.**  $d$  and  $T_g(H)$  for  $(\text{Ag}_{1-x}\text{Au}_x)_{0.9}\text{Mn}_{0.1}$ .

	$x$			
	0	0.007	0.03	0.05
$d$	0.095	0.129	0.203	0.252
$T_g$ (10 G) (K)	28.5	29.0	34.5	36.5
$T_g$ (2000 G) (K)	19.0	20.0	20.5	21.5

act in the chirality-driven extraordinary Hall effect mechanism by simultaneously measuring  $\rho_H$ ,  $M$  and the resistivity  $\rho$  for the series of AgMn alloys whose anisotropy is systematically controlled by doping with Au impurities.

The simultaneous measurement of  $\rho_H$ ,  $M$  and  $\rho$  were made from 8 to 50 K in a field of 2000 G under zero-field-cooled (ZFC) and field-cooled (FC) conditions. The samples used for the measurements are  $(\text{Ag}_{1-x}\text{Au}_x)_{0.9}\text{Mn}_{0.1}$  alloys with  $x = 0, 0.007, 0.03,$  and  $0.05$ . The details of the measurement and sample preparation are described in [1]. Table 1 shows the SG transition temperature  $T_g(H)$  and the anisotropy parameter  $d$  of the alloys. The anisotropy parameter  $d$  is defined as  $d \equiv D/J$ , where  $J$  and  $D$  are the exchange and anisotropy strengths respectively. The SG transition temperatures  $T_g(H)$  were determined from the magnetization measurements under ZFC and FC conditions. The values of  $d$  were calculated by using the formula in [6]. The value of  $T_g$  (10 G) increases in proportion to  $d^{0.8}$ , which is consistent with previous studies [6].

Figure 1(a) shows the temperature dependence of  $M$  for  $(\text{Ag}_{1-x}\text{Au}_x)_{0.9}\text{Mn}_{0.1}$  in a field of 2000 G [7]. The shift of  $T_g$  (2000 G) by doping Au impurities is also proportional to  $d^{0.8}$ . We observed that the doping has no effect on the magnitude of  $M$  in the high-temperature



**Figure 2.** Temperature dependence of  $R_s$  for  $(\text{Ag}_{1-x}\text{Au}_x)_{0.9}\text{Mn}_{0.1}$  in a field of 2000 G. It is noted that the temperature is divided by  $T_g$  (2000 G). The arrows mark  $T_g$  (2000 G).

region ( $T \geq 150$  K). On the other hand, the magnitude of  $M$  around  $T_g$  (2000 G) decreases with increasing concentration of Au impurities. This behaviour may be due to mean free path effects [8]. Figure 1(b) shows the temperature dependence of  $\rho_H$  for  $(\text{Ag}_{1-x}\text{Au}_x)_{0.9}\text{Mn}_{0.1}$  which was simultaneously measured with  $M$  [7]. The behaviour of  $\rho_H$  is similar to that of  $M$ , and the differences between ZFC and FC  $\rho_H$  appear below  $T_g$  (2000 G). The magnitude of  $\rho_H$  around  $T_g$  (2000 G) also decreases with Au concentration.

The Hall resistivity  $\rho_H$  is the sum of  $\rho_{\text{ord}}$  and  $\rho_{\text{ex}}$ . Extrapolations to high temperature to obtain an estimate of  $\rho_{\text{ord}}$  for the present samples indicate that  $R_0$  is about  $-8 \times 10^{-13} \Omega \text{ cm G}^{-1}$ .  $R_s$  is determined by using the value of  $R_0$  and the  $\rho_H$  and  $M$  data.

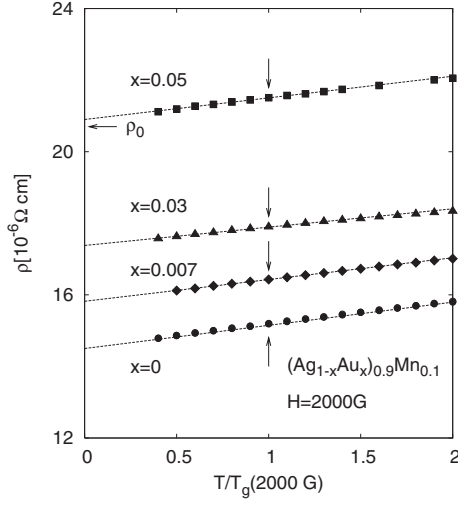
The temperature dependence of  $R_s$  for  $(\text{Ag}_{1-x}\text{Au}_x)_{0.9}\text{Mn}_{0.1}$  in a field of 2000 G is shown in figure 2. The differences  $\Delta R_s$  between the values of ZFC and FC  $R_s$  in the low-temperature region ( $T/T_g$  (2000 G)  $< 1$ ) are observed in all samples. The difference  $\Delta R_s$  clearly increases with the amount of Au impurities. In particular,  $\Delta R_s$  for  $(\text{Ag}_{0.95}\text{Au}_{0.05})_{0.9}\text{Mn}_{0.1}$  is as large as those of AuFe [1] and AuMn [2].

In the conventional theory [3],  $R_s = A\rho + B\rho^2$ , where  $\rho$  is the resistivity and  $A$  and  $B$  are constants relevant to the detailed band structure of the conduction electrons. The temperature dependence of  $\rho$ , as shown in figure 3, is monotonic even around  $T/T_g$  (2000 G) = 1, and the differences between ZFC and FC  $\rho$  are not observed in any samples. Therefore, the observed  $\Delta R_s$  are not explained by the conventional theory.

Tatara and Kawamura have shown that the uniform chirality  $\chi_0$  contributes to the extraordinary Hall resistivity  $\rho_{\text{ex}}$  by a perturbation expansion to the weak coupling s-d Hamiltonian [4] as follows:

$$\rho_{\text{ex}} = (A\rho + B\rho^2)M + C\chi_0, \quad (2)$$

where  $C$  is a constant relevant to the detailed band structure of the conduction electrons. The uniform chirality  $\chi_0$  is the sum of the local chirality  $\chi_{ijk} \equiv \mathbf{S}_i \cdot (\mathbf{S}_j \times \mathbf{S}_k)$  weighted by a geometrical factor which depends on the distance between the spins. Noting the geometrical factor, the contribution from  $\chi_{ijk}$  of three spins on the triangle  $P_i P_j P_k$  to  $\chi_0$  decays rapidly as  $\sim e^{-3r/2l}/(k_F r)^3$  [4], where  $r$  is the distance,  $k_F$  is the Fermi wave number,  $l$  is the mean free



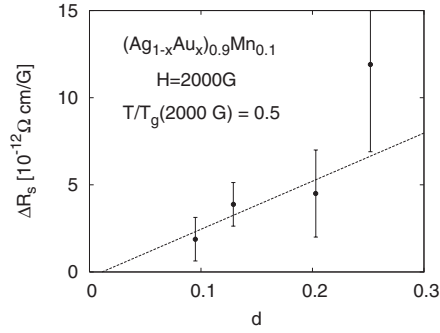
**Figure 3.** Temperature dependence of  $\rho$  for  $(\text{Ag}_{1-x}\text{Au}_x)_{0.9}\text{Mn}_{0.1}$  in a field of 2000 G. It is noted that the temperature is divided by  $T_g$  (2000 G). The arrows mark  $T_g$  (2000 G). The dotted lines show the result of the linear fitting below 30 K.  $\rho_0$  is the residual resistivity.

**Table 2.**  $\rho_0$  and  $l$  for  $(\text{Ag}_{1-x}\text{Au}_x)_{0.9}\text{Mn}_{0.1}$ .

	$x$			
	0	0.007	0.03	0.05
$\rho_0$ ( $10^{-6}$ $\Omega$ cm)	15	16	17	21
$l$ ( $\text{\AA}$ )	54	51	49	39

path and  $P_i$ ,  $P_j$  and  $P_k$  are the positions of  $S_i$ ,  $S_j$  and  $S_k$  respectively. This means that  $\chi_{ijk}$  of three spins on the triangle  $P_i P_j P_k$  having side-length up to  $l$  dominantly contributes to  $\chi_0$ . Therefore the contribution from  $\chi_{ijk}$  to the Hall effect disappears when the average distance  $r_{\text{ave}}$  between the spins is longer than  $l$ . Table 2 shows the residual resistivity  $\rho_0$  and  $l$  for  $(\text{Ag}_{1-x}\text{Au}_x)_{0.9}\text{Mn}_{0.1}$ . As shown in figure 3, values of  $\rho_0$  are determined from the extrapolation to  $T = 0$  in the temperature dependence of  $\rho$ . The mean free paths  $l$  are estimated by using the value of  $\rho_0$ , assuming that collisions of the conduction electrons with impurities dominate  $\rho$  in the low-temperature region. Because  $r_{\text{ave}} \simeq 6 \text{ \AA}$  in  $(\text{Ag}_{1-x}\text{Au}_x)_{0.9}\text{Mn}_{0.1}$ ,  $l$  is longer than  $r_{\text{ave}}$ . Therefore there are a lot of triangles  $P_i P_j P_k$  having side-length up to  $l$  and the contribution from  $\chi_{ijk}$  to  $\chi_0$  should appear in the present samples. However, since spins are frozen in a spatially random manner in the SG ordered state, the sign of  $\chi_{ijk}$  appears randomly, which inevitably leads to the vanishing of the uniform chirality,  $\chi_0 = 0$ .

It thus appears that the chirality-driven extraordinary Hall effect vanishes in bulk SG sample. To examine the possible coupling between  $\chi_0$  and  $M$ , Tatara and Kawamura have looked into the effective Hamiltonian of the spin-orbit interaction  $H_{\text{so}}$ , treating s-d interaction as a perturbation. They have also shown that the effective Hamiltonian which comes from the second-order contribution contains a term  $H_{\text{so}}^{(2)} \sim EM\chi_0$  [4], where  $E$  is a constant which represents the strength of the coupling between  $\chi_0$  and  $M$ . The chiral symmetry-breaking term  $H_{\text{so}}^{(2)}$  guarantees  $\chi_0$  to be induced if the sample is magnetized. This means that the  $M$  is a 'chiral field' conjugate to  $\chi_0$ . Then the chiral susceptibility  $X_\chi$  is defined as  $X_\chi \equiv \chi_0/EM$  and  $R_s$  is



**Figure 4.** Anisotropy parameter  $d$ -dependence of  $\Delta R_s$  for  $(\text{Ag}_{1-x}\text{Au}_x)_{0.9}\text{Mn}_{0.1}$  at  $T/T_g(2000 \text{ G}) = 0.5$ .

represented as follows [5]:

$$R_s \equiv \rho_{\text{ex}}/M = (A\rho + B\rho^2) + CEX_\chi. \quad (3)$$

Noting the chiral symmetry-breaking term  $H_{\text{so}}^{(2)}$  is essentially the DM anisotropy, two important implications of the DM anisotropy dependence of  $R_s$  are provided. First,  $\chi_0$  is not induced in the system with vanishing DM anisotropy even if the sample is magnetized. Therefore the chiral susceptibility term of  $R_s$  should vanish in the system with vanishing DM anisotropy,  $CEX_\chi = 0$ . Second, the strength of the coupling between  $\chi_0$  and  $M$  depends on the DM anisotropy. Therefore the chiral susceptibility term of  $R_s$  should depend on the DM anisotropy.

In our experimental results,  $\Delta R_s$ , which represents the term  $CE\Delta X_\chi$ , clearly increases with Au concentration, where  $\Delta X_\chi$  is the difference between the values of ZFC and FC  $X_\chi$ . Figure 4 shows the  $d$ -dependence of  $\Delta R_s$  for  $(\text{Ag}_{1-x}\text{Au}_x)_{0.9}\text{Mn}_{0.1}$  at  $T/T_g(2000 \text{ G}) = 0.5$ . One can see that  $\Delta R_s \sim 0$  when  $d = 0$  and  $\Delta R_s$  is roughly proportional to  $d$ . According to the theoretical prediction by Tataru and Kawamura [4], this observation indicates that the DM anisotropy acts as a ‘chiral symmetry-breaking field’ inducing  $\chi_0$  in the presence of  $M$ , which results in the chirality-driven extraordinary Hall effect in canonical SG alloys.

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