Evidence for Two-Dimensional Spin-Glass Ordering in Submonolayer Fe Films on Cleaved InAs Surfaces

Toshimitsu Mochizuki, Ryuichi Masutomi, and Tohru Okamoto

Department of Physics, University of Tokyo, 7-3-1, Hongo, Bunkyo-ku, Tokyo 113-0033, Japan

(Dated: October 24, 2008)

Magnetotransport measurements have been performed on two-dimensional electron gases formed at InAs(110) surfaces covered with a submonolayer of Fe. Hysteresis in the magnetoresistance, a difference in remanent magnetoresistance between zero-field-cooling procedures and field-cooling procedures, and logarithmic time-dependent relaxation after magnetic field sweep are clearly observed at 1.7 K for a coverage of 0.42 monolayer. These features are associated with spin-glass ordering in the Fe film.

PACS numbers: 75.70.Ak, 73.25.+i, 75.50.Lk

Spin glasses are magnetic systems with randomly competing interactions. They have attracted great interest during the last few decades [1, 2, 3]. Spin-glass models and related methods have also been useful in other areas of science such as simulation of protein folding [4] and optimization problems in computer science [5]. Most of the attempts to understand spin glasses have been concerned with the behavior in three dimensions. It is generally believed that, in two dimensions, the spin-glass ordering does not occur at nonzero temperature. The lower critical dimension of spin-glass ordering has been shown to be \( d_c > 2 \) for Heisenberg spins [6] and XY spins [7], and Ising spins with Gaussian distribution of disorder [8, 9]. Although the situation for the Ising model with bimodal \((\pm J)\) disorder was controversial [10, 11], recent theoretical investigations do not support the existence of the spin-glass phase for \( T > 0 \) [12, 13, 14]. On the other hand, numerical calculations have demonstrated that the spin-glass-like ordering temperature can be nonzero for a two-dimensional (2D) Ising model with random nearest-neighbor interactions and ferromagnetic second-neighbor interactions [15, 16]. Experimentally, spin-glass behavior was found in thin films [17, 18, 19, 20] and layered compounds [21, 22, 23]. However, no observation has been reported for a single layer system with strict two dimensionality.

Submonolayer films of magnetic materials adsorbed on nonmagnetic substrates are promising candidates for 2D spin-glass systems. A random distribution of adsorbates can be obtained by deposition at low substrate temperatures, where surface diffusion is minimal and island growth is limited. In the case that the sign of the interaction depends on the relative position of adatoms, a competition between ferromagnetic and antiferromagnetic interactions is expected. As the substrate, narrow band-gap III-V semiconductors have a remarkable property. It is well known that a two-dimensional electron gas (2DEG) can be easily formed on the surface of InAs and InSb. Photoelectron spectroscopy measurements have shown that the position of the Fermi level lies above the conduction-band minimum at cleaved (110) surfaces with various kinds of adsorbed materials [24, 25]. Recently the present authors have performed magnetotransport measurements on inversion layers formed on cleaved surfaces of \( p \)-type InAs [26, 27] and InSb [28] covered with submonolayers of Ag or alkali metals. The observed coverage dependence of the Hall mobility [26, 28] indicates that adatoms strongly affect electron scattering in the inversion layer. It seems feasible to probe the properties of adsorbed ultrathin films through transport measurements of adjacent conduction layers.

In this Letter, we report magnetotransport measurements of inversion layers formed on \textit{in-situ} cleaved InAs(110) surfaces, covered with a submonolayer Fe film at low temperatures. Hysteresis in the magnetoresistance is found in a narrow coverage range. At a coverage of 0.42 monolayer, the remanent magnetoresistance shows a clear difference between zero-field-cooling (ZFC) procedures and field-cooling (FC) procedures. It also exhibits a dependence on the direction of the applied magnetic field which corresponds to Ising-like anisotropy of the Fe film. A long-time relaxation behavior is observed after a magnetic field sweep. These results strongly indicate that the 2D spin-glass ordering occurs in the submonolayer Fe film.

The InAs samples used were cut from a Zn-doped single crystal with an acceptor concentration of \( 1.2 \times 10^{17} \text{cm}^{-3} \). Sample preparation and experimental procedures are similar to those used in Ref. [26]. Cleavage of InAs, subsequent deposition of Fe and transport measurements on the cleaved surface (3 mm × 0.4 mm) were performed at low temperatures in an ultrahigh vacuum chamber with a liquid \( ^4 \text{He} \) cryostat. The standard four-probe lock-in technique was used at 13.8 Hz with two current electrodes and four voltage electrodes prepared by deposition of gold films onto noncleaved surfaces at room temperature. The sample was mounted on a rotatory stage to control the magnetic field direction with respect to the surface normal. The electron density \( N_s \) and mobility \( \mu \) of the 2DEG were determined from the Hall measurements in a perpendicular magnetic field.

Figure 1(a) shows the magnetoresistance of the 2DEG...
In Fig. 1(c), we show a magnetoresistance curve at Θ = 0.17 ML where the hysteresis disappears. Positive B-dependence in the high-B region is also seen. Since similar positive magnetoresistance appears in 2DEGs covered with nonmagnetic materials, it should be attributed to intrinsic effects of 2DEGs, such as the orbital effect owing to the finite thickness of the inversion layer [21, 30], or the resistivity increase induced by the spin polarization [31, 32]. On the other hand, it is hard to explain the negative magnetoresistance observed in the low-B region for Θ = 0.42 ML [Fig. 1(a)] in terms of the characteristics of the 2DEGs. The Θ dependencies of $N_s$ and $\mu$ are gradual in the range of 0.08 ML $\leq$ Θ $\leq$ 0.50 ML, although the amplitude of the negative magnetoresistance changes drastically with Θ. We consider that the negative magnetoresistance and hysteresis are caused by changes in the magnetic state of the Fe film. Conduction electrons in the inversion layer move in a random potential induced by the spatial magnetization fluctuations of the Fe layer unless the exchange interaction is negligible. The negative magnetoresistance can be attributed to the suppression of the magnetization fluctuations with increasing average polarization. In contrast, an effect of the magnetization of conduction electrons on the magnetism of adsorbates is expected to be negligible since $N_s$ is 2 orders of magnitude smaller than the atomic density of adsorbates and the Pauli paramagnetic susceptibility is very small.

A recent calculation by Sacharow has shown that ferromagnetic structures are favorable in Fe[001] chains and antiferromagnetic structures are favorable in Fe[110] chains on InAs(110) [33]. Coexistence of ferromagnetic and antiferromagnetic interactions is expected since Fe adatoms in the present sample should be randomly distributed due to low substrate temperature deposition. The observed hysteresis in the magnetoresistance is associated with the irreversibility in a spin glass which appears at appropriate submonolayer coverages. By analogy with temperature-concentration phase diagrams for some three-dimensional systems [3], we think that the spin-glass phase exists between the ferromagnetic (higher-Θ) and paramagnetic (lower-Θ) phases. The reduction of $\rho$ during the initial magnetic field cycle cannot be related simply to the remanent magnetization since it is not recovered by applying a reverse magnetic field. The result suggests that a strong magnetic field has a persistent effect on the magnetization fluctuations in the spin-glass system. The remanent magnetoresistance can be removed only by annealing the sample. We found that the zero-magnetic-field resistivity returns to the initial value of 223.0 Ω after a thermal cycle up to 12 K or higher. From this we estimate a spin-glass transition temperature for Θ = 0.42 ML to be $T_g = 12$ K.

In general, the magnetic state of a spin-glass system depends strongly on the external magnetic field in which the sample was cooled from $T_g$. The difference in remanent...
magnetization between zero-field-cooled and field-cooled samples is observed in various materials \[3\]. In Fig. 2(a), the zero-magnetic-field resistivity \(\rho_0\) at \(T = 1.7\) K is shown for various cooling and magnetizing procedures. In ZFC procedures, the magnetic field excitation up to a maximum value \(B_{\text{max}}\) was performed at the measurement temperature of 1.7 K as illustrated in Fig. 2(b). On the other hand, it was performed at \(T_g = 12\) K in FC procedures [see Fig. 2(c)]. The cooling rate was 0.5 K/min in both procedures. For all series, \(\rho_0\) decreases as \(B_{\text{max}}\) increases and approaches a constant value in a high \(B_{\text{max}}\) regime where the spin polarization is expected to be completed. The observed \(B_{\text{max}}\)-dependence of \(\rho_0\) was roughly approximated by \(\rho_0(B_{\text{max}}) = \rho_0(0) - \Delta\rho/[1 + (B_0/B_{\text{max}})^2]\), where \(\rho_0(0) = 223.02\Omega, \Delta\rho = 1.58\Omega\) and \(B_0\) is a fitting parameter for each curve. The reduction of \(\rho_0(B_{\text{max}})\) for FC procedures is faster than that for ZFC procedures. This is consistent with a common feature of spin-glass systems—that FC magnetization is larger than ZFC magnetization \[3\]. The reduction of \(\rho_0\) depends also on the direction of the external magnetic field. The observed anisotropy suggests that the magnetic field strength required for the complete spin polarization is lowest in the direction perpendicular to the surface \((\varphi = 0^\circ)\) and the system has Ising-like (easy-axis) anisotropy.

A particularly interesting feature of spin glasses is the anomalously slow relaxation. The approximately logarithmic time dependence of the relaxation of the remanent magnetization has been reported for various spin-glass systems \[3\]. Figure 3(a) shows time evolution of \(\rho\) after magnetic field sweep at \(\varphi = 90^\circ\). (a) Data obtained at 4.0 K (upper curve) and 1.7 K (lower curve) after a sweep from 9 T to 0 T. (b) Data obtained at \(T = 1.7\) K and \(B = 1.5\) T after a down sweep from 2.0 T (upper curve) and an up sweep from 0.9 T (lower curve).
netization \[3, 36\] which can be interpreted as a delayed response to a magnetic field change. As discussed above, the resistivity of the 2DEG is considered as a probe of the magnetization fluctuations of the Fe layer. The observed $t$ dependence of $\rho$ suggests that the magnetization fluctuations always decrease when the spin-glass system relaxes toward a metastable state, irrespective of whether the average magnetization increases or decreases.

In summary, we have studied a magnetic state of the submonolayer Fe film through magnetotransport measurements of the 2DEG formed at the cleaved surface of InAs. Hysteresis behavior is observed in a narrow coverage range around $\Theta = 0.42$ ML. It is associated with the irreversibility of a spin-glass system of Fe adatoms. This interpretation is strongly supported by the observations of characteristic features of spin glasses. A clear difference between ZFC procedures and FC procedures is seen in the remanent magnetoresistance measurements and relaxation after magnetic field sweep exhibits a logarithmic time-dependence.

We thank T. Matsui for pointing out to us Ref. \[33\]. This work has partly supported by Grant-in-Aid for Scientific Research (B) (No. 18340080), Grant-in-Aid for Scientific Research on Priority Area “Physics of New Quantum Phases in Superclean Materials” (No. 20029005), and Grant-in-Aid for JSPS Foundation (No. 1811418) from MEXT, Japan.

[1] S. F. Edwards and P. W. Anderson, J. Phys. F 5, 965 (1975).
[2] D. Sherrington and S. Kirkpatrick, Phys. Rev. Lett. 35, 1792 (1975).
[3] K. Binder and A. P. Young, Rev. Mod. Phys. 58, 801 (1986).
[4] J. D. Bryngelson and P. G. Wolynes, Biopolymers 30, 177 (1990).
[5] S. Kirkpatrick, J. C. D. Gelatt, and M. P. Vecchi, Science 220, 671 (1983).
[6] J. R. Banavar and M. Cieplak, Phys. Rev. Lett. 48, 832 (1982).
[7] J. Maucourt and D. R. GREMPel, Phys. Rev. Lett. 80, 770 (1998).
[8] W. L. McMillan, Phys. Rev. B 30, 476 (1984).
[9] A. J. Bray and M. A. Moore, J. Phys. C 17, L463 (1984).
[10] T. Shirakura and F. Matsubara, Phys. Rev. Lett. 79, 2887 (1997).
[11] F. Matsubara, T. Shirakura, and M. Shiomi, Phys. Rev. B 58, R11821 (1998).
[12] N. Kawashima and H. Rieger, Europhys. Lett. 39, 85 (1997).
[13] A. K. Hartmann and A. P. Young, Phys. Rev. B 64, 180404(R) (2001).
[14] J. Houdayer, Eur. Phys. J. B 22, 479 (2001).
[15] N. Lemke and I. A. Campbell, Phys. Rev. Lett. 76, 4616 (1996).
[16] A. K. Hartmann and I. A. Campbell, Phys. Rev. B 63, 094423 (2001).
[17] L. Sandlund, P. Granberg, L. Lundgren, P. Nordblad, P. Svedlindh, J. A. Cowen, and G. G. Kenning, Phys. Rev. B 40, 869 (1989).
[18] L. Hoines, R. Stubi, R. Loloee, J. A. Cowen, and J. Bass, Phys. Rev. Lett. 66, 1224 (1991).
[19] P. Granberg, J. Mattsson, P. Nordblad, L. Lundgren, R. Stubi, J. Bass, D. L. Leslie-Pelecky, and J. A. Cowen, Phys. Rev. B 44, 4410 (1991).
[20] E. Morenzoni, H. Luetkens, T. Prokscha, A. Suter, S. Vongtragool, F. Galli, M. B. S. Hesselberth, N. Garifanov, and R. Khasanov, Phys. Rev. Lett. 100, 147205 (2008).
[21] A. Gavrin, J. R. Childress, C. L. Chien, Martinez, and M. B. Salamon, Phys. Rev. Lett. 64, 2438 (1990).
[22] T. Mori and H. Mamiya, Phys. Rev. B 68, 214422 (2003).
[23] R. Mathieu, J. P. He, Y. Kaneko, H. Yoshino, A. Asamitsu, and Y. Tokura, Phys. Rev. B 76, 014436 (2007).
[24] M. Getzlaff, M. Morgenstern, Chr. Meyer, R. Brochier, R. L. Johnson, and R. Wiesendanger, Phys. Rev. B 63, 205305 (2001), and references therein.
[25] M. G. Betti, V. Corradini, G. Bertoni, P. Casarini, C. Mariani, and A. Abramo, Phys. Rev. B 63, 155315 (2001).
[26] Y. Tsuji, T. Mochizuki, and T. Okamoto, Appl. Phys. Lett. 87, 062103 (2005).
[27] M. Minowa, R. Masutomi, T. Mochizuki, and T. Okamoto, Phys. Rev. B 77, 233301 (2008).
[28] R. Masutomi, M. Hio, T. Mochizuki, and T. Okamoto, Appl. Phys. Lett. 90, 202104 (2007).
[29] P. Štreda, P. Vašek, and M. Cukr, Phys. Rev. B 51, 11144 (1995).
[30] J. M. Heisz and E. Zaremba, Phys. Rev. B 53, 13594 (1996).
[31] T. Okamoto, K. Hosoya, S. Kawaji, and A. Yagi, Phys. Rev. Lett. 82, 3875 (1999).
[32] T. Okamoto, M. Ooya, K. Hosoya, and S. Kawaji, Phys. Rev. B 69, 041202(R) (2004).
[33] L. Sacharow, Ph. D. thesis, Institute of Applied Physics, University of Hamburg, 2006. http://www.physnet.uni-hamburg.de/services/fachinfo/search.php.
[34] Data for $t < 50$ sec are not shown because the temperature of the sample stage slightly fluctuated due to a technical problem of the temperature control system. For $t > 50$ sec, the fluctuations were not observed and we believe that the temperature difference between the sample and the resistance thermometer was negligible. After zero-field-cooling from 12 K to 1.7 K with 0.5 K/min, time dependence was not observed at least for $t > 50$ sec both in the sample resistivity and the thermometer resistance.
[35] For $\Theta = 0.33$ ML and 0.50 ML, the $t$-dependence of $\rho$ was found to be very weak after low-temperature magnetic field sweep.
[36] L. Lundgren, P. Svedlindh, P. Nordblad, and O. Beckman, Phys. Rev. Lett. 51, 911 (1983).