Anisotropic magnetoresistance components in (Ga,Mn)As

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Our experimental and theoretical study of the non-crystalline and crystalline components of the anisotropic magnetoresistance (AMR) in (Ga,Mn)As is aimed at exploring basic physical aspects of this relativistic transport effect. The non-crystalline AMR reflects anisotropic lifetimes of the holes due to polarized Mn impurities while the crystalline AMR is associated with valence band warping. We find that the sign of the non-crystalline AMR is determined by the form of spin-orbit coupling in the host band and by the relative strengths of the non-magnetic and magnetic contributions to the impurity potential. We develop experimental methods directly yielding the non-crystalline and crystalline AMR components which are then independently analyzed. We report the observation of an AMR dominated by a large uniaxial crystalline component and show that AMR can be modified by local strain relaxation. We discuss generic implications of our experimental and theoretical findings including predictions for non-crystalline AMR sign reversals in dilute moment systems.

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Anisotropic magnetoresistance (AMR) is a response of carriers in magnetic materials to changes of the magnetization orientation. Despite its importance in magnetic recording technologies the understanding of the microscopic physics of this spin-orbit (SO) coupling induced effect is relatively poor. Phenomenologically, AMR has a non-crystalline component, arising from the lower symmetry for a specific current direction, and crystalline components arising from the crystal symmetries [1,2]. In ferromagnetic metals, values for these coefficients can be obtained by numerical ab initio transport calculations [3], but these have no clear connection to the standard physical model of transport arising from spin dependent scattering of current carrying low mass s-states into heavy-mass d-states [4]. Experimentally, the non-crystalline and, the typically much weaker, crystalline AMR components in metals have been indirectly extracted from fitting the total AMR angular dependences [2].

Among the remarkable AMR features of (Ga,Mn)As ferromagnetic semiconductors are the opposite sign of the non-crystalline component (compared to most metal ferromagnets) and the crystalline terms reflecting the rich magnetocrystalline anisotropies [2,4,5,6,7,8,9,10,11,12]. Microscopic numerical simulations [6,12] consistently describe the sign and magnitudes of the non-crystalline AMR and capture the more subtle crystalline terms associated with e.g. growth-induced strain [8,12]. As in metals, however, the basic microscopic physics of the AMR still needs to be elucidated which is the aim of the work presented here.

Theoretically, we separate the non-crystalline and crystalline components by turning off and on band warping and match numerical microscopic simulations with model analytical results. This provides the physical interpretation of the origin of AMR, and of the sign of the non-crystalline term in particular. Experimentally, we obtain direct and independent access to the non-crystalline and crystalline AMR components using Hall bars fabricated along the principle crystalline axes and Corbino disk samples. The method is first established in the standard (Ga,Mn)As films before discussing the unique behavior we observe in ultra-thin low-conductive (Ga,Mn)As layers, which show a large, uniaxial crystalline component dominated AMR. Finally we demonstrate how crystalline AMR components can be strongly modified by local strain relaxation [12].

The experimental data presented in this paper were measured in compressively strained 25nm and 5nm Ga0.95Mn0.05As films grown by low temperature molecular beam epitaxy on GaAs [001] substrates. Optical lithography was used to fabricate Hall bars aligned along the [100], [010], [110] and [110] directions, of width 45μm with voltage probes separated by 285μm and Corbino disks of inner diameter 800μm and outer diameter 1400μm in which current flows radially in the plane of the material. Electron beam lithography was used to fabricate 1μm wide Hall bars in a 25nm Ga0.95Mn0.05 As film. All magnetoresistances were measured with the magnetization in the plane of the device, i.e., in the pure AMR geometry with zero (antisymmetric) Hall signal.

The phenomenological decomposition of the AMR of (Ga,Mn)As into various terms allowed by symmetry is obtained by extending the standard phenomenology [3], to systems with cubic [100] plus uniaxial [110] anisotropy. With this we write the longitudinal AMR as, $\Delta \rho_{xx}/\rho_{av} = C_I \cos 2\phi + C_U \cos 2\psi + C_C \cos 4\psi + C_{I,C} \cos (4\psi - 2\phi)$,
where $\Delta \rho_{xx} = \rho_{xx} - \rho_{av}$, $\rho_{av}$ is the $\rho_{xx}$ averaged over 360° in the plane of the film, $\phi$ is the angle between the magnetization unit vector $\hat{M}$ and the current $I$, and $\psi$ the angle between $\hat{M}$ and the [110] crystal direction. The four contributions are the non-crystalline term, the lowest order uniaxial and cubic crystalline terms, and a crossed non-crystalline/crystalline term. The purely crystalline terms are excluded by symmetry for the transverse AMR and we obtain, $\Delta \rho_{xy}/\rho_{av} = C_I \sin 2 \phi - C_{I,C} \sin(4 \psi - 2 \phi)$.

Microscopically we explain the emergence of the AMR components starting from the valence-band kinetic-exchange description of (Ga,Mn)As with metallic conductivities, which is an established qualitative and often semi-quantitative theoretical approach [14, 13]. The description is based on the canonical Schrieffer-Wolff transformation of the Anderson Hamiltonian which for (Ga,Mn)As replaces hybridization of Mn $d$-orbitals with As and Ga $sp$-orbitals by an effective spin-spin interaction of $L = 0, S = 5/2$ local-moments with host valence band states. These states, which carry all the SO-components, are the non-crystalline term, the lowest order between conductivity and lifetimes of carriers with the [110] crystal direction. The four contributions are the non-crystalline term, the lowest order between $\hat{M}$ and $I$, consistent with experiment. Our analysis also predicts that when the SO-coupling in the host band is of the form $s \perp k$, as in the Rashba-type 2D systems, or when Mn forms an isovalent pure magnetic impurity, e.g. in II-VI semiconductors, the sign of the non-crystalline AMR will be reversed.

In these dilute moment systems two distinct microscopic mechanisms lead to anisotropic carrier lifetimes, as illustrated in Fig. 1(a): One combines the SO-coupling in the carrier band with polarization of randomly distributed magnetic scatterers and the other with polarization of the carrier band itself resulting in an asymmetric band-spin-texture. Although acting simultaneously in real systems, theoretically we can turn both mechanisms on and off independently. We find that the former mechanism clearly dominates in (Ga,Mn)As which allows us to neglect spin-splitting of the valence band in the following qualitative discussion. This is further simplified by focusing on the non-crystalline AMR in the heavy-hole Fermi surfaces in the spherical, $s \parallel k$, spin-texture approximation [14] (see Fig. 1(a)) and considering scattering off a $\delta$-function potential $\propto (\alpha + \hat{M} \cdot s)$. Here $s$ and $k$ are the carrier spin-operator and wavevector, and $\alpha$ represents the ratio of non-magnetic and magnetic parts of the impurity potential. Assuming a proportionality between conductivity and lifetimes of carriers with $\mathbf{k} \parallel \mathbf{I}$ we obtain,

$$\frac{\sigma(\hat{M} \parallel I)}{\sigma(\hat{M} \perp I)} = \left(\alpha^2 + \frac{4}{12}\right) \left(\alpha^2 - \frac{4}{12}\right)^{-2}. \quad (1)$$

Therefore when $\alpha \ll 1$, one expects $\sigma(\hat{M} \parallel I) < \sigma(\hat{M} \perp I)$ (as is usually observed in metallic ferromagnets). But the sign of the non-crystalline AMR reverses at a relatively weak non-magnetic potential ($\alpha = 1/\sqrt{20}$ in the model), its magnitude is then maximized when the two terms are comparable ($\alpha = 1/2$), and, for this mechanism, it vanishes when the magnetic term is much weaker than the non-magnetic term ($\alpha \to \infty$).

Physically, carriers moving along $\hat{M}$, i.e. with $s$ parallel or antiparallel to $\hat{M}$, experience the strongest scattering potential among all Fermi surface states when $\alpha = 0$, giving $\sigma(\hat{M} \parallel I) < \sigma(\hat{M} \perp I)$. When the non-magnetic potential is present, however, it can more efficiently cancel the magnetic term for carriers moving along $\hat{M}$, and for relatively small $\alpha$ the sign of AMR flips.

Numerical simulations of hole scattering rates, illustrated in Fig. 1(a) on a color-coded minority heavy-hole Fermi surface, were obtained within the spherical

\begin{figure}[h]
\centering
\includegraphics[width=\textwidth]{fig1.png}
\caption{(a) Non-crystalline AMR in spherical bands: 2D Cartoons of AMR mechanisms and calculated anisotropic scattering rate on the 3D Fermi surface of the minority heavy-hole band in Ga$_{0.95}$Mn$_{0.05}$As. (b) Non-crystalline and crystalline AMR on warped bands: calculated anisotropic scattering rates for $\hat{M}$ \[100\] and \[110\] axes. (c) Calculated and (d) measured (at 4.2 K) longitudinal and transverse AMR for Ga$_{0.95}$Mn$_{0.05}$As as a function of the angle between $\hat{M}$ and $I$. The legend shows the direction of the current. The y-axes show $\Delta \rho/\rho_{av}$ shifted such that the minimum is at zero.}
\end{figure}
approximation but including the hole spin polarization, light-hole and split-off valence bands, and realistic non-magnetic and magnetic Mn impurity potentials [12]. The simulations confirm the qualitative validity of the analytical, non-crystalline AMR expressions of Eqs. (11). The additional crystalline AMR terms are obtained when the spherical approximation is relaxed and band warping is included in the Kohn-Luttinger Hamiltonian [13, 14].

The enhanced scattering of holes moving perpendicular to $\mathbf{M}$ seen in Fig.1(b) reflects the persistence of a strong non-crystalline component in the warped bands. The presence of the crystalline components in these anisotropic Fermi surfaces is also clearly apparent in Fig.1(b). In (Ga,Mn)As, the crystalline terms reflect the biaxial cubic anisotropy of these zincblende compounds combined with a [110] uniaxial component [15].

We conclude the theory discussion by showing in Fig.1(c) full numerical Boltzmann theory simulations of the AMR for a weakly (15%) compensated Ga$_{0.95}$Mn$_{0.05}$As material. The non-crystalline AMR contributes strongly and, as explained above, leads to a higher resistance state for $\mathbf{I} \perp \mathbf{M}$. Differences among AMRs for current along the [100], [110], and [110] directions show that cubic and uniaxial crystalline terms are also sizable. This phenomenology is systematically observed in experimental AMRs of weakly or moderately compensated metallic (Ga,Mn)As films. Typical data for such systems, represented by the 25nm Ga$_{0.95}$Mn$_{0.05}$As film with 3.6% AMR, are shown in Fig.1(d) for the Hall bars patterned along the [100], [010], [110], and [110] directions. In these measurements a saturating magnetic field of 1T is applied in the plane of the film and the magnetization vector follows the external field direction.

Above we have shown that diluted magnetic semiconductors like (Ga,Mn)As allow us to formulate the theory of AMR from the understanding of the very basic microscopic mechanisms. In what follows we focus on several unique experimental aspects of the AMR in these systems. The high crystalline quality and metallic character of the samples allow us to produce low contact resistant Hall bars accurately orientated along the principle crystallographic axes, from which it is possible to extract the independent contributions to the AMR. We are also able to fabricate low contact resistance Corbino disk samples for which the averaging over the radial current lines eliminates all effects originating from a specific direction of the current. Corbino measurements are possible in these materials because they are near perfect single crystals but with low carrier density and mobility (compared with single crystal metals) and so can have source-drain resistances large compared with the contact resistances.

Measured results for a Corbino device fabricated from the same 25nm Ga$_{0.95}$Mn$_{0.05}$As film as used for the Hall bars are shown in Fig. 2(a). The AMR signal is an order of magnitude weaker than in the Hall bars and is clearly composed of a uniaxial and a cubic contribution. Fig. 2(a) also shows the crystalline components of the AMR extracted by fitting the Hall bar data to the phenomenological longitudinal and transverse AMR expressions [20]. Fig. 2(b) shows the consistency for the coefficients $C_I, C_U$ and $C_C$ when extracted from the Hall bar and Corbino disk data over the whole range up to the Curie temperature (80K). Note that the uniaxial crystalline term, $C_U$, becomes the dominant term for $T \geq$30K. This correlates with the uniaxial component of the magnetic anisotropy which dominates for $T \geq$30K as observed by SQUID magnetometry measurements (not shown).

Our work shows that in (Ga,Mn)As ferromagnets, the symmetry breaking mechanism behind the previously reported [19] uniaxial magneto-crystalline anisotropy in the magnetization also contributes to the AMR.

We now discuss the unique AMR phenomenology observed on ultra thin (5nm) Ga$_{0.95}$Mn$_{0.05}$As films. Measurements on the Hall bars in Fig. 3(a) show that the AMR is very different from that observed in the 25nm film. Application of the phenomenological analysis to the Hall bar data shows that this behavior is a consequence of the crystalline terms dominating the AMR with the uniaxial component of the magnetic anisotropy being the largest. SQUID magnetometry on 5nm Ga$_{0.95}$Mn$_{0.05}$As films consistently shows that the uniaxial component of the magnetic anisotropy dominates over the whole temperature range [21]. The Corbino disk AMR data for a nominally identical 5nm film, shown in Fig. 3(b), confirm our observation of the highly unconventional 6% AMR totally dominated by the uniaxial crystalline term.

The 5nm films have lower Curie temperatures ($T_C \approx$30K) than the 25nm films and become highly resistive at low temperature indicating that they are close to the metal-insulator transition. The 25nm films show metallic behavior up to the lowest measured temperatures. The strength of the effect in the 5 nm films is remarkable and it is not captured by theory simulations assuming weakly disordered, fully delocalized (Ga,Mn)As valence bands. It might be related to the expectation that magnetic interactions become more anisotropic with...
increasing localization of the holes near their parent Mn ions as the metal-insulator transition is approached [14].

Finally, we demonstrate how the crystalline terms can be tuned by the use of lithographic patterning to induce an additional uniaxial anisotropy in very narrow Hall bars. In recent studies [13] it has been found that the pattern allows the in-plane compressive strain in the (Ga,Mn)As film to relax in the direction along the width of the Hall bar and this can lead to an additional uniaxial component in the magnetocrystalline anisotropy for bars with widths on the order of 1 µm or smaller. Figs. 3(c), (d) show the AMR of 45 µm wide bars and 1 µm wide bars fabricated from nominally identical 25 nm Ga0.95Mn0.05As wafers. For the 45 µm bars, the cubic crystalline symmetry leads to the AMR along [100] and [010] being larger than along [110] and [110]. For the narrow bars we observe the opposite relationship. This is consistent with the addition of an extra uniaxial component, whose presence in the magnetocrystalline anisotropy is confirmed by SQUID magnetometry measurements (not shown), which adds 0.8% to the AMR when current is along [110] and [110] and subtracts 0.4% when the current is along [100] and [010]. These post-growth lithography induced modifications are significant fractions of the total AMR of the parent (Ga,Mn)As material.

To conclude, we have described the non-crystalline AMR in (Ga,Mn)As as a combined effect of the SO-coupled spin-texture in the host band and polarized scatterers containing non-magnetic and magnetic impurity potentials. The additional crystalline terms are associated with band warping effects which reflect the underlying crystal symmetry. Our theory should apply to a large family of related dilute moment systems suggesting, e.g., that the non-crystalline AMR sign flips when the \( s \parallel k \) Kohn-Luttinger spin-texture is replaced by the \( s \perp k \) Rashba-type SO-coupling of asymmetric 2D systems, or when Mn forms an isovalent, pure magnetic impurity as is the case in II-VI semiconductor structures. On the experimental side we have established a technique for direct measurement of the crystalline AMR components by utilizing the Corbino disk geometry, which should also be applicable to other systems which combine high crystalline quality with relatively high resistivity. We report that in (Ga,Mn)As, ultra-thin films can be epitaxially grown with AMR dominated by a large uniaxial crystalline component and that in standard films the crystalline components can be modified by microscale lithography induced lattice relaxations.

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