Lateral diffusive spin transport in layered structures

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A one-dimensional (1D) theory of lateral spin-polarized transport is derived from the two-dimensional flow in the vertical cross section of a stack of ferromagnetic and paramagnetic layers. This takes into account the influence of the lead on the lateral current underneath, in contrast to the conventional 1D modeling by the collinear configuration of lead/channel/lead. Our theory is convenient and appropriate for the current in-plane configuration of an all-metallic spintronics structure as well as for the planar structure of a semiconductor with ferromagnetic contacts. For both systems we predict the optimal contact width for maximal magnetoresistance and propose an electrical measurement of the spin-diffusion length for a wide range of materials.

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Spintronics promises an increase in the logical expressibility of electronic circuits and the integration of nonvolatile magnetic memory.^{1,2} A quantitative and yet simple theory of spin transport is essential for interpreting experimental results and designing practical lateral devices which are imperative in large scale integrations. The most popular analytical approach to spin transport extends drift-diffusion equations to include the spin degree of freedom. $3-7$ Onedimensional (1D) equations of this kind have successfully explained the current-perpendicular-to-the-plane giant magnetoresistance⁴ and the spin injection from a ferromagnetic metal into a semiconductor in a vertical geometry.3,6,8–10 Theories of spin transport for the current in-plane focus on the boundary scattering to explain the magnetoresistive effect in ultrathin layered systems.11,12 However, many of the experiments involving nonmagnetic metals $13,14$ or semiconductors $15,16$ are performed in the planar geometry where the thickness of the layers exceeds the mean free path. For these systems, an application of either existing 1D models is a rough approximation, unable to account for some observed features whose interpretation required a numerical 3D treatment.17 The role of the contact width in lengthening the effective path between two leads versus the spin-diffusion length was mentioned.18 The effect of leakage from a two-dimensional electron gas into the ferromagnetic gate of finite lateral size on the spin currents in the channel has been investigated.¹⁹ What remains lacking is a simple theory in which the spin injection from contacts of finite width is normal to the spin current in the plane of the conduction channel.

In this paper, starting from a study of the two-dimensional current flow in the vertical cross section of the planar system, we derive an analytical 1D theory of spin transport in lateral structures. The analysis takes into account the lateral extent of the interface between ferromagnetic (FM) and normal (N) materials. We obtain a transparent physical picture and derive useful analytical formulas. The most important result obtained is the dependence of the magnetoresistance (MR) on the contact width. The knowledge of this aspect of lateral spin-valve physics is essential to the design of structures with high MR. In addition, our analysis suggests a technique to obtain the spin diffusion length of a paramagnetic material

by a simple set of electrical measurements. For compounds, such as silicon, that are inaccessible to optical characterization of spin polarization, $15,16$ development of the electrical method is of great importance.

Figure 1 depicts a typical planar system, homogeneous in the *z* direction. A model of a one-dimensional current flow from lead *B* via the normal region to lead *C* cannot describe the flow under the leads because of the simultaneous in-plane flow and the injection or extraction from above. Hence, we must first consider the two-dimensional transport governed by the spin-diffusion equation:

$$
\nabla^2 \mu_s(x, y) = \frac{\mu_s(x, y) - \mu_{-s}(x, y)}{(L_s)^2},
$$
 (1)

where μ_s is the spin-dependent electrochemical potential with $s=\pm$ denoting the spin species. The characteristic distance of the spin flip is $L_s = \sqrt{D_s \tau_{s-s}}$, where τ_{s-s} is the spinflip time and D_s is the diffusion constant of the spin component *s*. The measurable spin-diffusion length *L*, on whose scale the splitting of μ changes, is given by $L^{-2} = L_s^{-2} + L_{-s}^{-2}$. These equations hold for ferromagnetic and normal metals⁵ and also hold for nondegenerate semiconductors with the additional condition of a small electric field.⁶ Since the current is driven by the gradient of the electrochemical potential, the continuity of the normal component of the spin current across two adjacent layers *i* and *k* provides the boundary conditions,

FIG. 1. (Color online) A two-dimensional sketch of a planar system with ferromagnetic and normal materials. *A*–*D* represent boundaries which may be connected to the external circuit. All of the labeled lengths enter into our 1D transport equations.

$$
\sigma_s^i(\hat{n}^i \cdot \nabla \mu_s^i) = G_s^{i,k}(\mu_s^k - \mu_s^i) = -\sigma_s^k(\hat{n}^k \cdot \nabla \mu_s^k),\tag{2}
$$

where σ_s^i and \hat{n}^i refer to layer *i* conductivity and outward interface normal, respectively, and $G_s^{i,k}$ is the spin-dependent barrier conductance. Treating interfacial spin scattering requires additional relaxation terms in Eq. (2) as formally handled by Fert and Lee.²⁰ However, for a barrier which satisfies the condition in Eq. $(4a)$, this procedure²⁰ is equivalent to using Eq. (2) with renormalized barrier conductances whose spin selectivity $\tilde{G}^{i,k}_{+}/\tilde{G}^{i,k}_{-}$ is closer to unity. We will show that the optimal contact width is nearly independent on the latter.

Although Fig. 1 illustrates only two layers (normal and FM), the equations derived below hold for multilayers. To reduce the essential features of the 2D flow to 1D, we define the vertical (y) average of μ_s in each layer,

$$
\xi_s^i(x) = \frac{1}{h^i} \int_{y_0^i}^{y_1^i} dy \, \mu_s^i(x, y), \tag{3}
$$

with h^i being the thickness of the layer between its boundaries, (y_0^i, y_1^i) . For thin layers, $\mu_s^i(x, y)$ may be replaced by its vertical average $\xi_s^i(x)$. The conditions of validity follow the requirement that the gradient correction along y to μ_s in Eq. (3) be negligible to $O(h^2)$,

and

(a)
$$
h \ll \sigma_s / G_s
$$
,

$$
(b) \quad h < L_s,\tag{4}
$$

with the help of Eqs. (1) and (2) . Under these conditions we can derive a set of 1D equations governing the lateral (i.e., in the layer plane) spin transport. Integrating out the *y* dependence in Eq. (1) and using Eq. (2) yields

i

$$
\frac{\partial \xi_s^i(x)}{\partial x^2} = \frac{\xi_s^i(x) - \xi_{-s}^i(x)}{(L_s^i)^2} + \frac{G_s^{i,i+1}}{\sigma_s^i h^i} [\xi_s^i(x) - \xi_s^{i+1}(x)] + \frac{G_s^{i,i-1}}{\sigma_s^i h^i} [\xi_s^i(x) - \xi_s^{i-1}(x)].
$$
\n(5)

This kinetic theory assumes that the relevant length scales exceed the electron mean free path. Thus, the present analysis cannot address the quantum realm of the MR effect in ultrathin layered systems.¹¹

We now divide the lateral transport into regions of vertical stacks. Consider a region with a vertical stack of N_L layers. For example, in the region of width w_L covered by the left lead in Fig. 1, N_L =2, excluding the insulator. Transport in the layers of the stack is governed by $2N_L$ coupled members of Eq. (5) which in the matrix form are simply

$$
(\partial^2/\partial x^2)\xi = \mathbf{M} \cdot \xi,\tag{6}
$$

with the column vector ξ of elements ξ_s^i and the positive definite matrix M of elements from Eq. (5) . The solution has the form

$$
\xi(x) = \mathbf{1}(a + bx) + \sum_{j=1}^{N_L - 1} \mathbf{v}_j(p_j e^{\lambda_j x} + q_j e^{-\lambda_j x}), \tag{7}
$$

DERY, CYWIŃSKI, AND SHAM **PHYSICAL REVIEW B** 73, 041306(R) (2006)

where **1**, a column of ones, is an eigenvector of **M** with zero eigenvalue and **v**_j is an eigenvector with eigenvalue λ_{j}^2 . Interfacial spin scattering changes the values of \mathbf{v}_j and λ_j^2 but not the general form of the solution as the boundary conditions are linear²⁰ in the electrochemical potentials. The connection of the vertical stacks to the outside of the system and to each other is maintained by the boundary conditions. Between stacks they are given by the continuity of $\xi_s(x)$ and their first derivatives (currents) through each homogeneous layer. At the outermost boundaries (such as *A* and *D* in Fig. 1) the conditions are prescribed by the external driving terms of the electrochemical potential in the form of either a constant voltage maintained at an electrode or an injection current (including zero) through an interface. Applying currents and/or voltages to *B* and *C* interfaces result in the presence of inhomogenous terms in Eq. (6). These boundary conditions provide a unique set of solutions for the parameters a , b , p_j , q_j in Eq. (7). This method of a solution is a considerable simplification compared to solving the 2D spindiffusion equation for the same planar structure, and yet it includes the influence of the vertical layers on the lateral spin current flow. In the following, we will show the versatility of this simplified approach as well as testing its validity against the 2D solution.

The first illustrative example involves a semiconductor as the normal layer of the structure in Fig. 1. *A* and *D* are disconnected from the outside while *B* and *C* are voltage biased. The spin-diffusion length in the normal layer is L_N . We divide the normal channel into sections belonging to five vertical stacks as shown in Fig. 1. In the middle section of the channel, the solution is, $5,6$

$$
\xi_{\pm}(x) = (e/\sigma_N)Jx + \mu \pm (p_c e^{x/L_N} + q_c e^{-x/L_N}),
$$
 (8)

where *J*, μ , p_c , and q_c are constants to be determined by the boundary conditions. The total current flowing between the two leads is $I = Jh_N$ per unit length of the structure in the *z* direction. For the two sections outside the footprint defined by the two leads, we introduce the notion of the "open" versus "confined" geometry depending on whether $d_L, d_R \ge 0$ or $\ll L_N$. In both geometries the pattern of the net charge current occurs only between the leads *B* and *C*. This means that the outside sections lack the linear term of Eq. (8). Additionally the open geometry includes only an exponentially decaying solution away from the closer lead, and in contrast with the confined geometry, spin polarization extends noticeably outside of the footprint between *B* and *C*, thus reducing the spin accumulation densities. In each of the vertical stacks containing a lead, due to vastly different conductivities of the semiconductor and ferromagnetic metal we can decouple the equations for ξ in the *N* semiconductor and in the FM metal and solve for the eigenvalues of **M** in the former

$$
\lambda_{(s,c)}^2 = [\alpha + 1 \pm \sqrt{1 + \beta^2}]/(2L_N^2),
$$

$$
(\alpha, \beta) = 2L_N^2 (G_+ \pm G_-)/(\sigma_N h_N),
$$
 (9)

where the first of each pair of symbols (s, c) or (α, β) takes the upper sign. The electrochemical potential in the FM is

FIG. 2. (a) Magnetoresistance effect versus the contact width of a GaAs channel at 300 K. The solid (dashed) line denotes a confined (open) structure. The dots are the results of a 2D numerical computation in the confined geometry. (b) Magnetoresistive effect versus α for three cases of spin selectivity, β/α in the confined geometry, calculated for the optimal contact width corresponding to the same α value shown in (c).

practically a constant $(\xi^{FM} = -eV)$, and in the semiconductor layer under the biased lead is

$$
\xi_{\pm}(x) = -eV + (1 \pm \lambda)[pe^{\lambda_{s}x} + qe^{-\lambda_{s}x}]
$$

+ (\lambda \mp 1)[re^{\lambda_{c}x} + se^{-\lambda_{c}x}], (10)

where p , q , r , and s are constants to be determined by boundary conditions and $\lambda = \cot \left(\frac{1}{2} \tan^{-1} \beta \right)$. Consider the case that spin polarization is robust so that α and β are comparable. If $\alpha \ll 1$, then $\lambda_c \ll 1/L_N$ and $\lambda_s \sim 1/L_N$. The *s* mode is limited by the spin-diffusion constant, and it corresponds to spin accumulation ($\lambda \geq 1$ in this case). If $\alpha \geq 1$, then both eigenvalues are nearly independent of L_N , and neither of the eigenvectors is a pure spin mode $(\lambda \approx 1)$: the inhomogeneity of injection dominates the spatial dependence.

To illustrate the effect of the contact width on MR, we use the experimentally verified barrier parameters of a Fe/GaAs system at 300 K. 21,22 The tunneling conductances are of the order of $10^2 - 10^3 \Omega^{-1}$ cm⁻² with the ratio of spin-up to spindown conductance $G_{\perp}/G_{\perp} \simeq 2$. The *N* channel doping is n_0 $=4\times10^{15}$ cm⁻³ and the spin relaxation time is $\tau_s=80$ ps (Ref. 23), which corresponds to $L_N \approx 1 \mu m$. The thickness of the *N* channel is $h_N = 100$ nm, and the inner channel length is $d_m = 200$ nm.

The MR effect is defined as $(I^P - I^{AP})/I^P$ with $I^P(I^{AP})$ being the total current when the FM contacts are magnetized in parallel (antiparallel) directions. The calculated MR is shown in Fig. $2(a)$ as a function of the contact width, where for simplicity we have used $w=w_L=w_R$. In the confined geometry, $d_L = d_R = 0$, the effective 1D method (solid line) is compared with the results of 2D numerical calculation (dots), showing excellent agreement. We note the deleterious effect of the open geometry (dashed line) compared with the confined geometry (solid line) because of the weaker spin accumulation in the semiconductor channel which produces the MR effect.²⁴

(2006)

Note the existence of the optimal contact width for the maximum MR effect that arises out of the balance between spin injection and spin relaxation in the channel. The spin accumulation in the normal conduction channel is built up from injection along the width of the ferromagnetic contact. When the width is too small the resulting injection levels are not sufficient to evidently split the spin-dependent electrochemical potentials. Consequently the MR effect is small. On the other hand, when the contact width *w* exceeds the spin diffusion length L_N spin polarization diffuses in both directions beneath the wide contact and spin information from the far side is already "lost" when reaching the near side. In this case the build-up from vertical injection along the width of the contact beyond L_N becomes ineffective, the spin injection and the MR effect approach asymptotic values. The difference between the open and confined structures is removed, as can be seen from the merging of the solid and dashed lines in Fig. $2(a)$. The magnitude of MR at the optimal contact width depends also on the spin selectivity of the barriers, β in Eq. (9) , as illustrated in Fig. 2(b). Figure 2(c) shows a much weaker dependence on the spin selectivity of the optimal contact width in units of the spin-diffusion length (w_{opt}/L_N) . The ratio depends only on α as defined in Eq. (9). For α ≥ 1 , $w_{opt}/L_N \approx 2/\alpha$. For $\alpha \leq 1$, the optimal contact width w_{opt}/L_N is approximately $\frac{6}{5}$ or $\frac{3}{8}$ for the open or confined geometry, respectively. These results enable the extraction of the spin-diffusion length of a test material by measuring the MR of several structures with different contact widths from the same growth.

To demonstrate the use of our method for the all-metallic system, we turn to a second example. In the planar structure of Fig. 1, a constant current is injected from region *A* into a 10 nm thick paramagnetic metal layer. The structural parameters in terms of the spin-diffusion length of the *N* layer are $0.2d_L=0.2d_R=2d_M=L_N$. The FM layers *B* and *C* are of the same thickness and serve as floating contacts. We use the Al/Co system parameters at 300 K where the ratio of spin-up to spin-down conductance is $G_{+}/G_{-} = 2$ (Ref. 25) and their sum is $1.5 \times 10^9 \Omega^{-1}$ cm⁻². The normal layer intrinsic parameters are $\sigma_N = 3 \times 10^5 \Omega^{-1} \text{ cm}^{-1}$, $L_N = 0.35 \mu \text{m}$. In the Co layers the majority channel conductivity and spin-diffusion lengths are, respectively, $0.1\sigma_N$ and 12 nm. The respective values of the Co minority channel are $0.04\sigma_N$ and 20 nm. For the ease of toggling between *P* and *AP* configurations we use a width ratio of 2 between left and right Co regions. For the FM/ N regions the solutions are given by Eq. (7) . In the middle *N* channel the solution of $\xi_s(x)$ is given by Eq. (8) while at the outer channels the form of $\xi_s(x)$ lacks the diverging exponential.

For each magnetic configuration $(P \text{ or } AP)$ we calculate the voltage drop V_{BC} with the electrochemical potentials averaged along each Co layer. The MR is defined as 1 $-V_{BC}^{P}/V_{BC}^{AP}$. Figure 3(a) shows the MR effect. It peaks at the right contact width of $w_R \sim 0.5L_N$. The optimal contact width depends mostly on α , insensitive to varying the spin selectivity, the middle *N* channel length, or the outer *N* channel lengths (not shown). Figures $3(b)$ and $3(c)$ show, respectively, the current densities in the normal layer for parallel and antiparallel magnetic configuration at the optimal contact

FIG. 3. (a) The magnetoresistive effect versus the right contact width in the Al/Co system at 300 K. (b) The current density profile in the Al layer for a parallel magnetic configuration at the optimal contact width. Solid (dash) lines denote spin-up (down) current components. The total current is constant where in the dip regions along the Al/Co interfaces currents propagate in both layers. (c) Same as (b) for the antiparallel magnetic configuration.

width. The solid (dashed) lines depict spin-up and spin-down current components along the *x* direction. Despite the small MR effect, its observation is well within existing experimental abilities. Thus a similar analysis to the previous FM/

DERY, CYWIŃSKI, AND SHAM **PHYSICAL REVIEW B** 73, 041306(R) (2006)

semiconductor example leads to an electrical method to measure the spin-diffusion length of the normal material. However, the extraction of L_N requires that it greatly exceeds the FM spin diffusion length.

In summary, we have presented an effective 1D theory which describes the spin-dependent transport beneath and between ferromagnetic contacts in the lateral geometry. Within realistic constraints on the ratio between the layer conductivity and the contact conductance, the derived set of coupled linear equations governing the lateral diffusive spin currents is almost as accurate as the 2D spin diffusion equation but much simpler to use. This method retains the important role of the contact width, predicting an optimal contact width for the maximal magnetoresistive effect in the lateral spin valve. We have also quantified the effect of open channels, where spins can diffuse away from the active region underneath the FM contacts, in contrast with the confined structures, where the spin accumulation is kept between the two contacts. Our results provide crucial guidelines for design of lateral spin-valve type devices. They also yield a method of extracting the spin-diffusion length in an allelectrical measurement. At room temperature where the spindiffusion length lies in the submicron region, this method does not have the difficulty of wavelength resolution in the optical techniques.

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