

2012

Spin injection from a half-metal at finite temperatures

Kirill D. Belashchenko

University of Nebraska-Lincoln, belashchenko@unl.edu

James K. Glasbrenner

University of Nebraska-Lincoln, james.glasbrenner.ctr@nrl.navy.mil

Aleksander L. Wysocki

University of Nebraska-Lincoln, awysocki@iastate.edu

Follow this and additional works at: <http://digitalcommons.unl.edu/physicsbelashchenko>

Belashchenko, Kirill D.; Glasbrenner, James K.; and Wysocki, Aleksander L., "Spin injection from a half-metal at finite temperatures" (2012). *Kirill Belashchenko Publications*. 20.

<http://digitalcommons.unl.edu/physicsbelashchenko/20>

This Article is brought to you for free and open access by the Research Papers in Physics and Astronomy at DigitalCommons@University of Nebraska - Lincoln. It has been accepted for inclusion in Kirill Belashchenko Publications by an authorized administrator of DigitalCommons@University of Nebraska - Lincoln.

Spin injection from a half-metal at finite temperatures

K. D. Belashchenko,^{*} J. K. Glasbrenner, and A. L. Wysocki*Department of Physics and Astronomy and Nebraska Center for Materials and Nanoscience, University of Nebraska-Lincoln, Lincoln, Nebraska 68588, USA*

(Received 14 September 2012; published 4 December 2012)

Spin injection from a half-metallic electrode in the presence of thermal spin disorder is analyzed using a combination of random matrix theory, spin-diffusion theory, and explicit simulations for the tight-binding s - d model. It is shown that efficient spin injection from a half-metal is possible as long as the effective resistance of the normal metal does not exceed a characteristic value, which does not depend on the resistance of the half-metallic electrode but, rather, is controlled by spin-flip scattering at the interface. This condition can be formulated as $\alpha \lesssim l/l_{sf}^N T_c^{-1}$, where α is the relative deviation of the magnetization from saturation, l and l_{sf}^N are the mean-free path and the spin-diffusion length in the nonmagnetic channel, and T_c is the transparency of the tunnel barrier at the interface (if present). The general conclusions are confirmed by tight-binding s - d model calculations. A rough estimate suggests that efficient spin injection from true half-metallic ferromagnets into silicon or copper may be possible at room temperature across a transparent interface.

DOI: 10.1103/PhysRevB.86.224402

PACS number(s): 72.25.Hg, 72.25.Mk, 75.47.—m, 85.75.—d

I. INTRODUCTION

Many spintronic devices depend on the injection, manipulation, and detection of spin-polarized currents in semiconductors or normal metals.^{1–3} Spin injection can also be utilized as a tool to probe the spectroscopic properties of strongly correlated and spin-orbit-coupled systems.^{1,4} Thus, understanding the mechanisms of spin injection is of interest for a variety of fundamental and practical applications. Basic theory of spin injection across an F/N (ferromagnet/normal metal) interface in the linear response regime was worked out by Johnson and Silsbee.⁵ The spin polarization of the injected current may be conveniently expressed as^{1,6}

$$P_j = \frac{P_\sigma r_F + P_\Sigma r_c}{r_F + r_N + r_c}, \quad (1)$$

where $P_\sigma = (\sigma_\uparrow - \sigma_\downarrow)/(\sigma_\uparrow + \sigma_\downarrow)$, σ_\uparrow and σ_\downarrow are the spin-resolved conductivities of the ferromagnetic electrode, P_Σ is defined similarly to P_σ for the spin-dependent interface conductance, and r_F and r_N are the effective resistances of the ferromagnet and normal metal, respectively. The effective interface resistance is denoted r_c . For the ferromagnet $r_F = (\sigma_\uparrow + \sigma_\downarrow)l_{sf}^F/(4\sigma_\uparrow\sigma_\downarrow)$ and for the normal metal $r_N = l_{sf}^N/\sigma_N$, where l_{sf}^F and l_{sf}^N are the spin-diffusion lengths in the ferromagnet and in the normal metal. The quantities r_F and r_N are called the effective resistances. Expression (1) is valid under the assumptions of the two-current model,⁷ i.e., when the spin-diffusion lengths are much longer than the mean-free paths.⁸ Note that nonlinear effects in bipolar semiconducting junctions^{9,10} cannot be described within the linear-response theory and are beyond the scope of the present consideration.

Spin injection from a ferromagnet into a semiconductor is subject to the so-called conductivity mismatch problem.¹¹ For a typical choice of materials we have $l_{sf}^N \gg l_{sf}^F$ and $\sigma_N \ll \sigma_\uparrow\sigma_\downarrow/(\sigma_\uparrow + \sigma_\downarrow)$. This implies that $r_F \ll r_N$, and if the interface resistance r_c is also low compared to r_N , the injected current is unpolarized, $P_j \ll 1$. In order to circumvent this problem, one can introduce a highly resistive, spin-selective barrier at the interface, such as a naturally occurring Schottky barrier or an artificially inserted tunnel junction.⁶ According to

Eq. (1), a large r_c (comparable to or greater than r_N) combined with an appreciable P_Σ results in a finite P_j . Efficient spin injection into GaAs and Si from transition-metal electrodes was successfully achieved based on this principle.^{12–15}

The situation can be visualized with the help of an effective resistor circuit, such that in each spin channel s the ferromagnet and the normal metal have resistances l_{sf}^F/σ_s and $2l_{sf}^N/\sigma_N$, respectively, and the two spin channels are connected in parallel.¹⁶ This effective circuit correctly reproduces both the spin polarization of the current near the interface and the resistance of the junction in excess of what would be measured if the interface were replaced by a node in the circuit.¹⁶

Half-metallic ferromagnets¹⁷ are conducting in one spin channel and insulating in the other, which makes them attractive candidates as electrode materials for spintronic devices.¹ The situation at zero temperature is simple, as there is only one conducting channel (“spin up”), and the injected current should be fully spin-polarized. Many materials, particularly among Heusler compounds, have been theoretically predicted using band structure calculations to be half-metallic,¹⁸ although reliable experimental confirmation is often complicated by surface effects.¹⁹ High magnetoresistance values were achieved in magnetic tunnel junctions^{20–23} and spin valves²⁴ with epitaxial Co-based Heusler-alloy electrodes. Large nonlocal spin signals, 10 times higher compared to those of conventional electrodes, were also demonstrated in lateral spin valves with transparent ohmic interfaces.^{25–27} In all of these experiments the spin signal is considerably reduced at room temperature but remains appreciable. Further, Ramsteiner *et al.*²⁸ demonstrated spin injection from Co₂FeSi into an (Al,Ga)As light-emitting diode structure with an efficiency of at least 50%. Based on their device design, they concluded that the Schottky barrier cannot be present at the interface and argued that the observed efficient spin injection “casts doubt onto the common belief that tunneling is a prerequisite for spin injection from a metal into a semiconductor.”

So far the theoretical analysis of spintronic devices with half-metallic electrodes has been based^{25–27} on the standard spin-diffusion model,^{1,5,6,8,29} which assumes the existence of

two weakly coupled conducting channels in each material.^{7,8} However, this validity criterion are not satisfied in true half-metals, and our present goal is to develop an appropriate formalism for such devices, which should include the effects of thermal spin fluctuations at finite temperatures. Due to these fluctuations, the electron wave functions lose their pure spin character, and the density of states (DOS) in a half-metal acquires nonzero projection onto the “spin-down” channel, which is gapped at zero temperature. However, this state can be viewed as a small perturbation of the fully collinear spin state by fluctuating transverse magnetic fields, so that the local spin direction for all electronic eigenstates is fluctuating within a narrow cone around the magnetization direction. In other words, the number of eigenstates is not doubled, but rather they acquire a small spin-down component. In this situation one cannot apply the two-current model, in which independent distribution functions are introduced for spin-up and spin-down electrons, and the concept of the spin-diffusion length also becomes meaningless. Therefore, Eq. (1) cannot be directly applied to spin injection from a half-metallic electrode at $T \neq 0$.

In the following, we analyze the spin injection from a half-metallic electrode in the linear response regime but without making the assumptions of the two-current model leading to Eq. (1). We start with general considerations in Sec. II and then proceed to analyze the elastic scattering region using the random matrix theory in Sec. III. Here we derive the formula for spin-injection efficiency, which is similar to (1) but with the effective resistance \tilde{r}_F being controlled by spin-flip scattering probabilities at the interface. A generalized statement of the conductivity mismatch follows from the unitarity of the scattering matrix. Based on these results, we then describe the half-metallic spin-injection system within the spin-diffusion theory in Sec. IV. The formula for spin-injection efficiency is generalized in a natural way to the case of a finite spin-diffusion length in the normal region. In Sec. V we discuss the behavior of spin-injection efficiency, and, finally, in Sec. VI we support our conclusions with explicit tight-binding s - d model calculations. The conclusions are summarized in Sec. VII.

II. A HALF-METAL AT A FINITE TEMPERATURE

The electronic structure of a half-metal at $T = 0$ has a band gap in one of the spin channels. If we now consider a thermal fluctuation resulting in a small canting of individual local spin moments, we can imagine, on the level of the self-consistent field theory in the localized basis, that the effective fields at different atomic sites have been rigidly rotated off of the magnetization axis by small angles. (The tight-binding representation is assumed for simplicity and is not essential for the physical argument.) This is a common approach to spin fluctuations within the noncollinear density functional theory, whereby the spin moments are assumed to fluctuate adiabatically slowly compared with the electron hopping times.³⁰ This approximation is justified by the fact that typical times associated with magnon dynamics are much longer than the electron momentum relaxation time. The Hamiltonian of a system with such adiabatic spin fluctuation

can then be represented as³¹

$$H\{\hat{n}_i\} = \sum_i U(\hat{n}_i) H_i U^\dagger(\hat{n}_i) + K, \quad (2)$$

where \hat{n}_i is the unit vector parallel to the spin moment at site i , H_i is the on-site contribution to the Hamiltonian from site i , and K is the spin-diagonal kinetic (hopping) part. We can now make a unitary transformation to the new “rotated” local basis in which H_i is diagonal, which is effected by unitary matrices $U(\hat{n}_i)$. In this new basis the Hamiltonian is

$$\tilde{H}\{\hat{n}_i\} = \sum_i H_i + \sum_{ij} U^\dagger(\hat{n}_i) K_{ij} U(\hat{n}_j). \quad (3)$$

At zero temperature there are only states of a particular spin (say, “up”) near the Fermi level. Deviation of the unitary matrices in (3) from unity at $T \neq 0$ introduces hybridization between local spin-up and spin-down states, as well as some randomness in the spin-conserving hopping matrix elements. Bloch states near the Fermi level acquire a small admixture of spin-down character, and the DOS in the global basis acquires a spin-down component, but no new Bloch states appear near the Fermi level. This means that the bulk of a half-metal at finite (but low) temperatures can be treated as having one effective spin channel. This situation is qualitatively different from that of a conventional ferromagnet with two independent spin channels, even if they have very different resistivities. While the conventional ferromagnet has two independent occupation functions and chemical potentials for the two spin channels, a half-metal has only one. Transport across an interface with a normal metal is discussed in the subsequent sections.

III. SPIN INJECTION IN THE SCATTERING FORMALISM

A spin-injection device can be analyzed by treating the F/N interface as an elastic scattering region embedded between diffusive regions and by matching the solution of the scattering problem with the solution of the spin-diffusion equation. For a conventional two-channel ferromagnetic electrode the well-known result is given by Eq. (1). However, as we argued in Sec. II, the two-current model is inapplicable for a half-metallic electrode. The purpose of this section is to clarify the role of spin coherence in spin injection from a half-metal at finite temperatures, i.e., in the presence of spin disorder.

Since the half-metal, as argued above, has only one effective spin channel, there is no analog of the spin-diffusion length for it. Therefore, inelastic scattering should not affect the properties of spin injection, and we may treat the whole half-metallic electrode as an elastic scatterer. As we will see in the next section, matching with the solution of the spin-diffusion equation in the normal metal should simply replace the resistance of the normal region with its effective resistance $r_N = \rho_N l_{sf}^N$. Therefore, we first consider the entire spin-injection device disregarding inelastic scattering altogether.

An elastic spin-injection device can be considered in the formalism of the scattering theory. We assume that the F/N device is connected on both sides to equilibrium reservoirs via ideal Landauer leads. Apart from these leads we introduce an auxiliary lead in the N region at a distance from the interface (a few mean-free paths) such that the quantum interference effects occurring at the interface are left entirely on the left-hand

side of this lead. Each of the two regions can be described by a scattering matrix,

$$\hat{S}_F = \begin{pmatrix} \hat{r}_1 & \hat{t}'_1 \\ \hat{t}_1 & \hat{r}'_1 \end{pmatrix}, \quad \hat{S}_N = \begin{pmatrix} \hat{r}_2 & \hat{t}'_2 \\ \hat{t}_2 & \hat{r}'_2 \end{pmatrix}, \quad (4)$$

where, in the standard way, the matrix \hat{t}_1 contains amplitudes for transmission from the conducting channels of the left electrode across the F into the conducting channels of the fictitious lead, and similarly for the other sub-blocks. At this point we allow the F region to have an arbitrary magnetic configuration. The transmission matrix \hat{t} of the entire F/N junction is

$$\hat{t} = \hat{t}_2(1 - \hat{r}'_1\hat{r}_2)^{-1}\hat{t}_1. \quad (5)$$

The charge and spin currents flowing across the junction are proportional, respectively, to C and C_s :

$$C = \text{Tr} \hat{t}\hat{t}^+, \quad C_s = \text{Tr} \hat{\sigma}_z \hat{t}\hat{t}^+, \quad (6)$$

and we are interested in the spin polarization $P_j = C_s/C$. Since we are considering the junction as an elastic scattering region, the spin current in the normal region is conserved. There is no loss of generality from singling out the z axis, because its direction is unspecified.

Following the approach of Waintal *et al.*,³² we now introduce the polar decomposition³³ of the matrix \hat{S}_N ,

$$\hat{S}_N = \begin{pmatrix} \hat{U} & 0 \\ 0 & \hat{V}' \end{pmatrix} \begin{pmatrix} \sqrt{1-T} & i\sqrt{T} \\ i\sqrt{T} & \sqrt{1-T} \end{pmatrix} \begin{pmatrix} \hat{U}' & 0 \\ 0 & \hat{V} \end{pmatrix}, \quad (7)$$

where T is the matrix of the eigenvalues of $\hat{t}_2\hat{t}_2^+$, while \hat{U} , \hat{U}' , \hat{V} , and \hat{V}' are unitary matrices, which are all diagonal in spin space. Since the fictitious node can be introduced at a sufficient distance from the surface to eliminate all quantum interference effects, we can safely use the isotropic approximation,³⁴ i.e., assume that the spatial factors of the unitary matrices \hat{U} , \hat{U}' , \hat{V} , and \hat{V}' are distributed uniformly in the unitary group. We substitute (5) into (6), use (7) for \hat{t}_2 and \hat{r}'_2 , and integrate over the unitary ensemble. This integration is easily performed using the method in Ref. 35 to the leading order in the number of conducting channels in the leads. In this leading order, each unitary matrix is matched to its own conjugate, resulting in a ladder diagram.³⁵ Averaging over the eigenvalues of $\hat{t}_2\hat{t}_2^+$ is performed simultaneously. The result can be written in the form

$$C_s = \sum_{\lambda\mu} \sigma_\lambda^z [1 - \hat{R}(1 - T_N)]_{\lambda\mu}^{-1} T_\mu T_N. \quad (8)$$

Here λ and μ denote a pair of spin indices, σ_λ^z is the $\hat{\sigma}_z$ matrix written as a vector (1,0,0, -1), T_N is the probability of transmission through the N region, \hat{R} is a 4×4 matrix³² with elements

$$R_{\lambda\mu} \equiv R_{\sigma\sigma',ss'} = \frac{1}{N_{\text{ch}}} \sum_{mn} (r'_1)_{m\sigma,ns} (r'_1)_{m\sigma',ns'}^*, \quad (9)$$

where m and n enumerate the N_{ch} conducting channels in the auxiliary lead, and T_μ is a 4-vector with elements

$$T_\mu \equiv T_{ss'} = \frac{1}{N_{\text{ch}}} \sum_{mn\sigma} (t_1)_{ms,n\sigma} (t_1)_{ms',n\sigma}^*. \quad (10)$$

In this last expression $n\sigma$ labels the channels of the lead feeding the F region. The expression for C is obtained from (8) by replacing σ_λ^z with a 4-vector representation of the unit matrix $1_\lambda = (1,0,0,1)$. Note that $T_\mu = \sum_\lambda T_{\mu\lambda} 1_\lambda$, where $T_{\mu\lambda}$ is defined as $R_{\mu\lambda}$ but with matrix elements of t_1 instead of r'_1 .³⁶

The unitarity of \hat{S}_F requires that $\hat{t}_1\hat{t}_1^+ + \hat{r}'_1\hat{r}'_1^+ = 1$. This condition implies that $T_\lambda = \sum_\mu (\delta_{\lambda\mu} - R_{\lambda\mu}) 1_\mu$. Substituting this into (8) we find that C_s vanishes to first order in T_N . Since the charge current is proportional to T_N , this leads to $P_j \rightarrow 0$ at $T_N \rightarrow 0$. In the limit of a two-channel device with weak coupling between the channels, this result reduces to the conductivity mismatch obstacle for spin injection.¹¹ However, our result is more general, because it is valid for any magnetic structure of the F region and for any choice of the z axis. The only exception is the case of a half-metal at $T = 0$ with no spin-flip scattering at the interface, for which the only nonzero component of T_μ is $T_{\uparrow\uparrow}$ (in the reference frame where the z axis is aligned with the magnetization). In this exceptional case we obviously have $P_j = 1$ at any T_N .

Let us now find the spin polarization P_j for a finite T_N , assuming that the electrode is an axially symmetric (i.e., collinear) magnet with spin disorder. For a macroscopic interface, the summation over the conducting channels automatically averages $R_{\lambda\mu}$ and T_μ in (8) over the spin-disorder ensemble. This self-averaging does not necessarily occur in a point contact, in which case an additional averaging over the spin-disorder configuration is required for the spin current, (8), and its charge counterpart. Re-expanding the inverse matrix in (8), we can obtain a series of terms describing multiple scatterings at the interface. Since the electron scatters repeatedly from the same spin-disorder configuration, the averages of the matrix products do not decouple. However, since correlations between successive scattering events do not change the asymptotic behavior of P_j , it is a reasonable approximation to replace \hat{R} and T_μ with their averages $\langle \hat{R} \rangle$ and $\langle T_\mu \rangle$ even for a point contact.

Let us assume that spin-orbit coupling at the surface is weak and that all spin-flip processes are dominated by spin-disorder scattering. Then the matrices $\langle \hat{R} \rangle$ and $\langle T_\mu \rangle$ should be invariant with respect to rotation in spin space around the magnetization axis. This condition implies that $\langle T_{\uparrow\downarrow} \rangle$ and $\langle T_{\downarrow\uparrow} \rangle$ vanish, along with all elements $\langle R_{\sigma\sigma',ss'} \rangle$ with $\sigma - \sigma' \neq s - s'$. The $\langle \hat{R} \rangle$ matrix is thus block diagonal. From the structure of (8) it is clear that we are only interested in the 2×2 block spanned by indices 1 and 4. (The 22 and 33 diagonal elements represent the spin-mixing conductance,³⁷ which turns out to be irrelevant to the problem at hand.) As seen from (9), the diagonal elements of this block are the total spin-conserving reflection probabilities for spin-up and spin-down electrons, $R_\uparrow = \langle R_{\uparrow\uparrow,\uparrow\uparrow} \rangle$ and $R_\downarrow = \langle R_{\downarrow\downarrow,\downarrow\downarrow} \rangle$, while the off-diagonal elements are the total spin-flip reflection probabilities $R_{\uparrow\downarrow} = \langle R_{\uparrow\downarrow,\downarrow\downarrow} \rangle$. Reciprocity requires that $\langle R_{\uparrow\downarrow,\downarrow\downarrow} \rangle = \langle R_{\downarrow\downarrow,\uparrow\uparrow} \rangle$. Let us also denote $T_\uparrow = T_{\uparrow\uparrow}$ and $T_\downarrow = T_{\downarrow\downarrow}$. (Note that T_\uparrow and T_\downarrow include both spin-conserving and spin-flip processes.) We can now calculate the spin polarization from (8):

$$P_j = \frac{P_i \tilde{r}_F}{\tilde{r}_F + r_N}, \quad (11)$$

where $P_t = (T_\uparrow - T_\downarrow)/(T_\uparrow + T_\downarrow)$, $r_N G_N = (1 - T_N)/(2T_N)$,

$$\frac{1}{4G_N \tilde{r}_F} = \frac{T_\uparrow T_\downarrow}{T_\uparrow + T_\downarrow} + R_{\uparrow\downarrow}, \quad (12)$$

and $G_N = (e^2/h)N_{\text{ch}}$. Note that P_t is the spin polarization of the current injected into the auxiliary lead if the N region is detached from it.

Expression (11) includes the effects of spin disorder but not the effects of inelastic spin relaxation. For a conventional (not half-metallic) electrode, spin relaxation must be included on both sides of the junction. In the presence of spin-flip processes at the interface, the solution of the spin-diffusion equations becomes rather complicated³⁸ even if the spin-flip reflection $R_{\uparrow\downarrow}$ is neglected. The situation is simpler in the case of a half-metallic electrode, because inelastic spin relaxation should only be included in the normal region. In the next section we will see that in this case the elastic resistance of the normal region r_N in (11) should simply be replaced with its effective resistance.

Note that interfacial spin-flip scattering due to spin-orbit interaction has been studied in some detail for metallic N/N and F/N interfaces.^{39,40} Temperature-dependent interfacial spin-flip scattering in the presence of nonequilibrium spin accumulation was suggested as a source of asymmetric response in a nonlocal spin valve.⁴¹

IV. SEMICLASSICAL THEORY

In the previous section we found that under rather general assumptions the scattering at the interface between the half-metal and the normal metal is described completely by spin-dependent transmission probabilities and the spin-flip reflection probability on the normal-metal side. The effects of spin coherence are effectively eliminated by spin-disorder averaging. We can therefore use the standard semiclassical treatment, taking into account that the half-metal has only one spin channel and incorporating spin-flip scattering at the interface. Apart from giving a complementary picture of spin injection, this treatment confirms the expectation about the role of the spin-diffusion length in the normal metal and shows the invariance of the results with respect to the location of the left lead.

Instead of treating the whole F/N device as an elastic scatterer, we now consider only the interfacial F/N region (a few mean-free paths on both sides) embedded between infinite diffusive regions. The half-metallic (F) region carries only one spin channel (even at finite temperature), but the N region has two channels. Similarly to Rashba's treatment of the F/N junction with spin-flip transmission at the interface,³⁸ the interface is assigned the spin-flip conductance $\Sigma_{\uparrow\downarrow}$ in addition to the spin-conserving $\Sigma_{\uparrow\uparrow}$. In addition to these terms, we also need to introduce spin relaxation in the normal metal due to spin-flip scattering at the interface. Physically, even if the F electrode is insulating, the spin accumulation in the normal metal can relax through interfacial spin-flip scattering. Introducing the appropriate electrochemical potential drops at the interface, the spin-dependent currents on the normal-metal side of the interface can be written as

$$j_\uparrow^N(0) = \Sigma_{\uparrow\uparrow}(\zeta_\uparrow^N - \zeta^F) + \tilde{R}_{\uparrow\downarrow}(\zeta_\uparrow^N - \zeta_\downarrow^N), \quad (13)$$

$$j_\downarrow^N(0) = \Sigma_{\uparrow\downarrow}(\zeta_\downarrow^N - \zeta^F) + \tilde{R}_{\uparrow\downarrow}(\zeta_\downarrow^N - \zeta_\uparrow^N), \quad (14)$$

where the new term is the one with $\tilde{R}_{\uparrow\downarrow}$. Matching with the solution of the spin-diffusion equation can be worked out in the usual way.³⁸ The terms containing bulk conductivity in the F region drop out, and after some algebra we reproduce Eq. (11) with r_N now being the effective resistance $\rho_N I_{sf}^N$ (as anticipated), P_t replaced by $P_\Sigma = (\Sigma_{\uparrow\uparrow} - \Sigma_{\uparrow\downarrow})/(\Sigma_{\uparrow\uparrow} + \Sigma_{\uparrow\downarrow})$, and

$$\frac{1}{4\tilde{r}_F} = \frac{\Sigma_{\uparrow\uparrow}\Sigma_{\uparrow\downarrow}}{\Sigma_{\uparrow\uparrow} + \Sigma_{\uparrow\downarrow}} + \tilde{R}_{\uparrow\downarrow}. \quad (15)$$

Equation (15) is equivalent to (12) with the replacement $G_N T_\uparrow \rightarrow \Sigma_{\uparrow\uparrow}$, $G_N T_\downarrow \rightarrow \Sigma_{\uparrow\downarrow}$, and $G_N R_{\uparrow\downarrow} \rightarrow \tilde{R}_{\uparrow\downarrow}$. At first sight, there is a discrepancy, because $\Sigma_{\uparrow\uparrow}$ and $\Sigma_{\uparrow\downarrow}$ are the interface conductances, while T_s are the total transmission probabilities of the entire half-metallic electrode. However, these expressions are, in fact, consistent, because \tilde{r}_F is invariant with respect to the choice of the boundary of the interface region at which the chemical potential ζ_F is evaluated. In order to see this, let us rewrite Eqs. (13) and (14) for the same F/N junction with a different choice of this boundary and denote the new chemical potential (at that boundary) ζ_0^F . This can be viewed as a simple redefinition of the thickness of the interface region. The chemical potentials on the normal side of the interface, however, are evaluated at the same point. The conductance and reflectance parameters corresponding to the new choice of ζ_0^F are denoted $\Sigma_{\uparrow\uparrow}^0$, $\Sigma_{\uparrow\downarrow}^0$, and $\tilde{R}_{\uparrow\downarrow}^0$.

For a half-metallic electrode the spin polarization of the current injected into the N region under the condition $\zeta_\uparrow^N = \zeta_\downarrow^N$ is determined by the ratio $\alpha = \Sigma_{\uparrow\downarrow}/\Sigma_{\uparrow\uparrow}$, which should depend only on temperature. Therefore, $\Sigma_{\uparrow\downarrow}/\Sigma_{\uparrow\uparrow} = \Sigma_{\uparrow\downarrow}^0/\Sigma_{\uparrow\uparrow}^0$. The charge current is

$$j = \Sigma_{\uparrow\uparrow}(\xi_\uparrow + \alpha\xi_\downarrow) = \Sigma_F \Delta\zeta_F, \quad (16)$$

where we denote $\xi_s = \zeta_s^N - \zeta^F$ and $\Delta\zeta_F = \zeta^F - \zeta_0^F$, and Σ_F is the conductance of the half-metallic region between the points where ζ^F and ζ_0^F are evaluated.

Equating the two different expressions for the same spin-dependent currents, (13) and (14), we can write

$$(\Sigma^0 - \Sigma)\xi_\uparrow + \Sigma^0\Delta\zeta_F + (\tilde{R}^0 - \tilde{R})(\xi_\uparrow - \xi_\downarrow) = 0, \quad (17)$$

$$\alpha(\Sigma^0 - \Sigma)\xi_\downarrow + \alpha\Sigma^0\Delta\zeta_F - (\tilde{R}^0 - \tilde{R})(\xi_\uparrow - \xi_\downarrow) = 0, \quad (18)$$

where we have simplified the notation by dropping indices: $\Sigma = \Sigma_{\uparrow\uparrow}$, $\tilde{R} = \tilde{R}_{\uparrow\downarrow}$, and similarly for Σ^0 and \tilde{R}^0 . Furthermore, substituting $\Delta\zeta_F$ from (16), we obtain a system of two linear homogeneous equations for ξ_\uparrow and ξ_\downarrow . For this system to have a solution, the determinant of the coefficient matrix should vanish, which leads to

$$\frac{\alpha}{1+\alpha}(\Sigma^0 - \Sigma) + \tilde{R}_0 - \tilde{R} = 0. \quad (19)$$

This expression implies that \tilde{r}_F defined in (15) does not depend on the definition of the boundary of the interface region on the half-metallic side. Physically, this property follows from the unitarity of the scattering matrix and cannot be satisfied without introducing the spin-flip reflection terms in (13) and (14).

Spin injection from a half-metallic electrode may be schematically represented by the equivalent resistor circuit

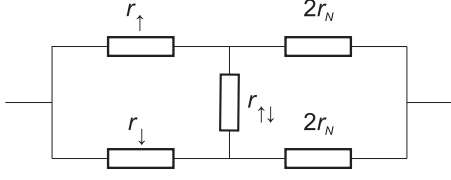


FIG. 1. Equivalent resistor circuit for spin injection from a half-metal.

shown in Fig. 1. The resistances are defined as $r_{\uparrow\downarrow} = 1/\tilde{R}_{\uparrow\downarrow}^\infty$, $r_\uparrow = 1/\Sigma_{\uparrow\uparrow}$, and $r_\downarrow = 1/\Sigma_{\uparrow\downarrow}$. Interestingly, the effective resistance $4\tilde{r}_F$ in (15) can be measured between the terminals of $r_{\uparrow\downarrow}$ if the normal-metal part of the circuit is disconnected and the left terminal is left open. P_Σ is given by $(r_\downarrow - r_\uparrow)/(r_\downarrow + r_\uparrow)$. The physical location of the left terminal of the circuit can be selected anywhere inside the half-metal. According to the arguments presented above, the change of this location redefines the three resistances r_\uparrow , r_\downarrow , and $r_{\uparrow\downarrow}$ while leaving $r_\uparrow/r_\downarrow \sim \alpha$, \tilde{r}_F , and P_j invariant. Note that the degradation of magnetic order at the interface may significantly affect P_Σ and \tilde{r}_F .

V. SPIN-INJECTION EFFICIENCY FOR A HALF-METALLIC ELECTRODE

The conductance and reflectance parameters defined in Eqs. (13) and (14) depend only on the properties of the interface region, and not on the properties of the bulk half-metallic region attached to it. Therefore, the spin-injection efficiency does not depend on the thickness of the half-metallic electrode. This result is valid as long as the half-metallic region is not so thin as to violate the assumptions of the diffusion theory. In practice, this means that it should be thick compared to the electronic mean-free path.

Using this property, we can formally include an arbitrarily thick half-metallic layer in the definition of the interfacial region, so that the conductances $\Sigma_{\uparrow\uparrow}$ and $\Sigma_{\uparrow\downarrow}$ are made very low (and thus the resistances r_\uparrow and r_\downarrow in Fig. 1 very high). Then the first term on the right-hand side of (15) is negligible compared to the corresponding asymptotic value $\tilde{R}_{\uparrow\downarrow}^\infty$, which is always finite at non-zero temperature. Thus, efficient spin injection is possible only when $\tilde{R}_{\uparrow\downarrow}^\infty \lesssim r_N^{-1}$. In other words, spin injection is suppressed at $r_N \gg (\tilde{R}_{\uparrow\downarrow}^\infty)^{-1} = r_{\uparrow\downarrow}^\infty$. On one hand, this condition is similar to the conductivity mismatch, because it sets a certain limit for r_N . On the other hand, the physical picture is quite different, because this limit is not related to the conductivity of the half-metal.

For generality let us allow for the existence of a magnetically unpolarized resistive barrier (such as a tunnel junction) at the interface, and let T_c denote the transmission probability across this barrier. From the first term in (15) we can deduce $\tilde{r}_F \sim (\alpha G_N T_c)^{-1}$. Alternatively, we can find $\tilde{R}_{\uparrow\downarrow}^\infty$ by adding a large resistor on the left of T_c with transmission probability $T_F \ll T_c$. In order to reflect with a spin flip, an electron incident from the normal-metal side must first tunnel across the barrier in order to reach the spin-disordered region; this gives a factor T_c . Since we require $T_F/T_c \rightarrow 0$, the electron is reinjected back into the N region with probability 1. The probability of spin flip adds a factor α , so we obtain $\tilde{R}_{\uparrow\downarrow}^\infty \sim \alpha G_N T_c$ and

confirm the above result for \tilde{r}_F . An electron can also scatter with a spin flip on the transverse exchange field introduced by the spin density penetrating across the barrier. This mechanism contributes in the same order to $\tilde{R}_{\uparrow\downarrow}^\infty$.

In the ohmic regime (low-resistance interface with $T_c \sim 1$), we have $G_N \tilde{r}_F \sim 1/\alpha$. Since $G_N r_N \sim l_{sf}^N/l$, where l is the mean-free path in the normal metal, we find that spin disorder suppresses spin injection when $\alpha \gg l/l_{sf}^N$. At small T we expect $\alpha \approx \langle \theta^2 \rangle / 4$, where θ is the polar angle of the injected spinor. The parameter α is approximately related to the reduced magnetization $m = M(T)/M(0)$ of the half-metal as $2\alpha \approx 1 - m$. (This quantity is proportional to the partial minority-spin DOS in the global spin basis.) Thus, the above condition shows the range of temperatures for which ohmic spin injection from a half-metal may be possible.

It is interesting to compare this result with the case of a two-channel ferromagnet with the same spin polarization of the DOS, for which the efficiency of ohmic spin injection is given by Eq. (1) with $r_c = 0$. Setting $\rho_\downarrow \sim \rho_\uparrow/\alpha$, we find $r_F \sim \rho_\uparrow l_{sf}^F/\alpha$, which should be compared to $\tilde{r}_F \sim \rho_N l/\alpha$ in the case of a half-metallic electrode. The dependence on the spin polarization of the globally defined DOS is similar, but the overall factor is different: the product $\rho_\uparrow l_{sf}^F$ is replaced with $\rho_N l$ in the case of a half-metal. For spin injection in semiconductors from metals, typically $\rho_N \gg \rho_\uparrow$, while l_{sf}^F is usually fairly small.³⁹ Thus, the effective resistance of a half-metallic electrode may be much higher compared to that of a conventional ferromagnet with a similar spin polarization of the DOS, which is an advantage for practical applications. On the other hand, since \tilde{r}_F does not depend on the resistivity of the half-metal, there is no benefit in increasing this resistivity. In particular, a magnetic semiconductor should not necessarily be a better spin injector than a highly conductive half-metal, assuming that half-metallicity is maintained at the interface in both cases.

For an actual device (e.g., F/N/F) it is necessary that l_{sf}^N is not small compared to the length L of the channel, and l/l_{sf}^N should be replaced with l/L if $l_{sf}^N \gg L$. Thus, for ohmic spin injection from a half-metal it may be beneficial to use a lightly doped semiconducting channel in order to maximize the mean-free path there. This is contrary to the conventional conductance mismatch considerations, according to which it is desirable to decrease ρ_N by increasing the doping concentration. The mean-free path in lightly doped silicon may be as high as 30 nm at room temperature.⁴² For a short channel with $L \sim 300$ nm this would allow ohmic spin injection at $\alpha \lesssim 0.1$. Although this is an order-of-magnitude estimate, at face value it allows spin injection for $M(T)/M(0) \gtrsim 0.8$. In elemental transition metals the reduced magnetization drops to 0.8 at about 75% of the Curie temperature, so this limitation is not very restrictive, particularly since the half-metallic gap may be closed by magnetic disorder at much lower temperatures.⁴³ A similar estimate applies to nonlocal spin valves with a copper channel, where at room temperature the mean-free path is of the order of 30 nm, and the spin-diffusion length is a few hundred nanometers.³⁹ It is possible that efficient spin injection across a transparent interface observed in Ref. 28 can be understood in a similar way.

From the point of view of interface engineering, it is always necessary to avoid the depletion region near the

surface.¹⁵ On the other hand, we would like to point out that in the ohmic regime the existence of interface states in the “wrong” spin channel does not necessarily preclude spin injection, as it would with a tunnel barrier. If these states are strongly hybridized with the normal region, they can be regarded as a part of the corresponding spin channel. (See Ref. 44 for a related discussion.) It is, however, important that the magnetic continuity and ordering at the interface are maintained. Otherwise, partially ordered regions or “loose spins” can provide strong spin-flip scattering at relatively low temperatures, thereby violating the $\alpha \lesssim l/l_{sf}^N$ inequality and suppressing spin injection. Thus, interface design based on chemical similarity of the F and N regions⁴⁵ may in practice be counterproductive, because it may be expected to facilitate interdiffusion.

With a tunnel barrier at the interface, the temperature range allowing efficient spin injection extends to $\alpha \lesssim (l/l_{sf}^N)/T_c$. As in the case of a conventional ferromagnetic electrode, the tunnel barrier is favorable for spin injection. In the case $T_c \ll l/l_{sf}^N$ spin injection is possible at any temperature, and its efficiency is proportional to $P_I = (1 - \alpha)/(1 + \alpha)$. (Of course, this assumes that the half-metal continues to behave as a single-channel conductor at elevated temperatures; real materials with a small half-metallic gap do not necessarily behave in this way.)

VI. TIGHT-BINDING CALCULATIONS

In this section we verify the conclusions of the general theory using tight-binding calculations for a specific realization of a half-metal based on the s - d model. Static spin disorder is introduced by randomizing the directions of the exchange fields on different sites according to the mean-field distribution function corresponding to the specified value of the magnetization. The spin-injection device is treated as an elastic system. We use a single-band Hamiltonian with nearest-neighbor interactions in the simple cubic lattice. For the half-metallic region, the energies in one spin channel are made very high in order to lift it far above the Fermi level. This is the limit of a large s - d exchange integral. The half-metallic and normal regions are sandwiched between two nonmagnetic leads. The hopping and band center parameters are selected as shown schematically in Fig. 2. Calculations were performed for supercells with a 10×10 cross section. The 2×2 conductance matrix $G_{ss'}$ was obtained using the Landauer-Büttiker approach, averaging over 100 configurations of spin and Anderson disorder. Brillouin-zone integration was performed using a 5×5 mesh, which was sufficient for convergence. To simulate diffusive transport, random disorder was applied to both half-metallic and semiconductor regions. Figure 2 illustrates the relative amplitude of random disorder relative to the band widths.

Since the spin current in the normal region is conserved, the spin polarization of the current flowing into the right electrode is identified with the spin-injection efficiency:

$$P_G = \frac{G_{\uparrow\uparrow} + G_{\downarrow\uparrow} - G_{\uparrow\downarrow} - G_{\downarrow\downarrow}}{G_{\uparrow\uparrow} + G_{\downarrow\uparrow} + G_{\uparrow\downarrow} + G_{\downarrow\downarrow}}. \quad (20)$$

According to the results in the previous sections, the dependence of P_G on the total resistance of the normal layer reflects

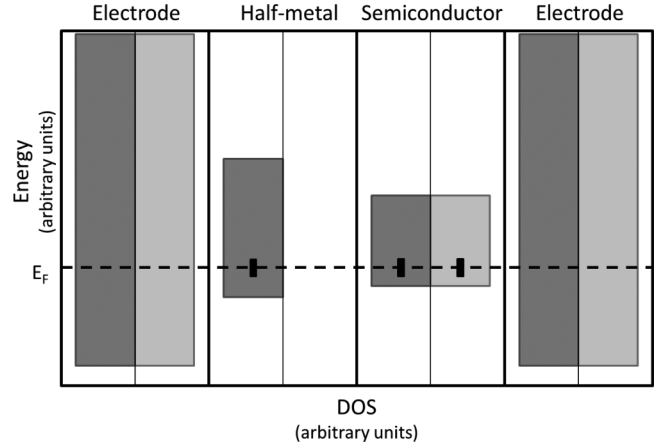


FIG. 2. Schematic of the band alignment for the spin-injection device without spin disorder. Darker and lighter bands correspond to majority and minority spins. The minority-spin band in the half-metallic region is shifted up beyond the energy range shown here. The dashed horizontal line shows the Fermi level, and the vertical black bars show the amplitudes of random disorder.

the dependence of spin-injection efficiency on the effective resistance r_N .

In the following we verify the following properties of the spin-injection efficiency P_G for a half-metallic electrode: (i) the independence of P_G from the thickness of the half-metal; (ii) the form, (11), of the dependence of P_G on the resistance of the normal region; (iii) the dependence of \tilde{r}_F on the magnetization of the half-metal, $\tilde{r}_F \sim \alpha^{-1}$; and (iv) the dependence of \tilde{r}_F on the transparency of a thin tunnel barrier inserted at the interface, $\tilde{r}_F \sim T_c^{-1}$.

The inset in Fig. 3(a) shows P_G as a function of the thickness of the half-metallic region at $m = M/M(0) = 0.9$, with a 50-monolayer-thick normal region. It is shown that P_G

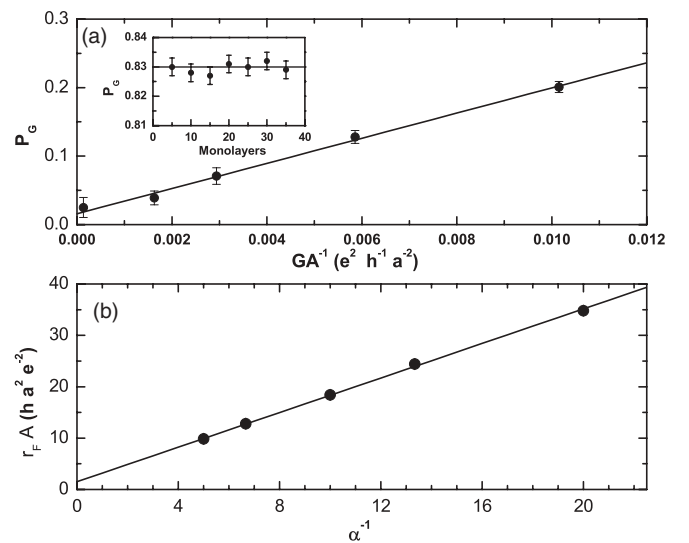


FIG. 3. (a) Conductance polarization as a function of the inverse resistance-area product for the reduced magnetization $m = 0.8$. Inset: The same quantity as a function of the thickness of the half-metal at $m = 0.9$. (b) Effective resistance \tilde{r}_F as a function of α^{-1} , where $\alpha = (1 - m)/2$.

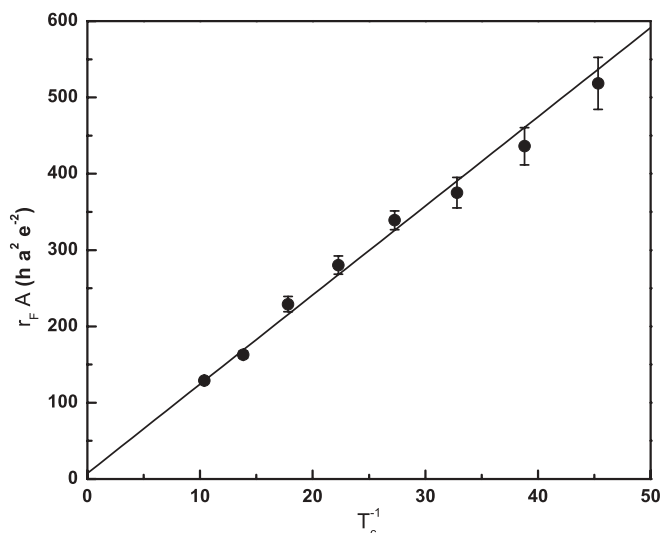


FIG. 4. Effective resistance \tilde{r}_F as a function of T_c^{-1} for fixed magnetization $m = 0.6$.

is independent of the thickness of the half-metal, in agreement with the general results. In all subsequent calculations the thickness of the half-metal is fixed at five monolayers.

In order to obtain the asymptotic dependence of P_G on the resistance of the normal region, we have added a tunnel barrier of variable height and thickness between the semiconductor and the right lead. This is necessary because otherwise the localization effects become important when the disordered normal region is made too long,⁴⁶ and the diffusive scaling breaks down. The length of the normal region is fixed at 50 monolayers. Figure 3(a) shows the dependence of P_G on the total resistance of the scattering region (which is dominated by the auxiliary tunnel barrier) for the reduced magnetization of $m = M/M(0) = 0.8$. It is clearly shown that P_G goes to 0 linearly with r_N^{-1} , in agreement with Eq. (11).

Next we evaluate the dependence of the effective resistance \tilde{r}_F on the magnetization m of the half-metal. To this end, for a given m the P_G is calculated from the configurationally averaged spin-dependence conductances for a set of thicknesses of the normal region ranging from 10 to 200 monolayers (all within the range where weak localization effects are undetectable). The $P_G(r_N)$ dependence is then fitted to Eq. (11) for each m . The magnitude of P_i decreases with decreasing magnetization but always remains somewhat larger than m . (This is likely due to the fact that the conduction electrons sample spin disorder over a few sites, effectively decreasing the transverse fields.) The dependence of \tilde{r}_F on the magnetization

is shown in Fig. 3(b), where $\alpha = (1 - m)/2$, as above. The linear dependence $\tilde{r}_F \propto \alpha^{-1}$ confirms the predictions of the general model, in which $\tilde{r}_F \propto R_{\uparrow\downarrow}^{-1} \sim \alpha^{-1}$. This divergence of \tilde{r}_F at low temperatures may be used experimentally as a signature of a true single-channel half-metal, although at very low temperatures the interfacial spin-orbit scattering may take over and cut off the divergence.

Finally, we considered the effect of a single-monolayer tunnel barrier at the F/N interface. We set $m = 0.6$ for the half-metal and varied the thickness of the semiconductor region from 10 to 200 monolayers, as above. As above, the P_G calculated from the averaged spin-dependent conductances was fitted to Eq. (11), extracting the $\tilde{r}_F(T_c)$ dependence. T_c was varied by changing the height of the tunnel barrier. Figure 4 shows the results supporting the inverse relationship $\tilde{r}_F \propto T_c^{-1}$. Together with the results shown in Fig. 3, we find $\tilde{r}_F \propto (\alpha T_c)^{-1}$, as expected.

VII. CONCLUSIONS

We have analyzed the spin injection from a half-metallic electrode into a normal (or semiconducting) region in the presence of thermal spin disorder. The two-current model with independent populations of the two spin channels is inapplicable to a half-metallic ferromagnet. The spin-injection efficiency is described by Eq. (11), in which r_N is the conventional effective resistance of the normal metal, while \tilde{r}_F is controlled by spin-flip scattering at the interface with the ferromagnet. Although \tilde{r}_F does not depend on the thickness of the half-metallic layer, its dependence on the spin polarization of the density of states and on the contact resistance is similar to the case of a conventional ferromagnet. Explicit simulations for the tight-binding s - d model confirm these general conclusions. In the case of a transparent interface, efficient spin injection is possible in the temperature range corresponding to $\alpha \lesssim l/l_{sf}^N$, where α is the relative deviation of the (surface) magnetization from saturation, and l is the mean-free path in the N region. A rough estimate suggests that efficient spin injection from half-metallic Co-based Heusler alloys into silicon or copper may be possible at room temperature across a transparent interface. Adding a tunnel barrier at the interface with transparency T_c extends this temperature range to $\alpha \lesssim l/l_{sf}^N T_c^{-1}$.

ACKNOWLEDGMENTS

We are grateful to Andre Petukhov, Evgeny Tsymbal, and Igor Žutić for useful discussions. This work was supported by NSF Grant No. DMR-1005642 and by Research Corporation through a Cottrell Scholar Award (K.B.).

*belashchenko@unl.edu

¹I. Žutić, J. Fabian, and S. Das Sarma, *Rev. Mod. Phys.* **76**, 323 (2004).

²M. Johnson, in *Handbook of Spin Transport and Magnetism*, edited by E. Y. Tsymbal and I. Žutić (CRC Press, Boca Raton, FL, 2012), p. 115.

³R. Jansen, *Nat. Mater.* **11**, 400 (2012).

⁴L. W. Cheuk, A. T. Sommer, Z. Hadzibabic, T. Yefsah, W. S. Bakr, and M. W. Zwierlein, *Phys. Rev. Lett.* **109**, 095302 (2012).

⁵M. Johnson and R. H. Silsbee, *Phys. Rev. B* **35**, 4959 (1987).

⁶E. I. Rashba, *Phys. Rev. B* **62**, R16267 (2000).

⁷N. F. Mott, *Proc. R. Soc. A* **153**, 699 (1936).

- ⁸T. Valet and A. Fert, *Phys. Rev. B* **48**, 7099 (1993).
- ⁹I. Žutić, J. Fabian, and S. Das Sarma, *Phys. Rev. Lett.* **88**, 066603 (2002).
- ¹⁰I. Žutić, J. Fabian, and S. C. Erwin, *J. Phys.: Condens. Matter* **19**, 165219 (2007).
- ¹¹G. Schmidt, D. Ferrand, L. W. Molenkamp, A. T. Filip, and B. J. van Wees, *Phys. Rev. B* **62**, R4790 (2000).
- ¹²A. T. Hanbicki, B. T. Jonker, G. Itkos, G. Kioseoglou, and A. Petrou, *Appl. Phys. Lett.* **80**, 1240 (2002).
- ¹³S. A. Crooker, M. Furis, X. Lou, C. Adelman, D. L. Smith, C. J. Palmstrom, and P. A. Crowell, *Science* **309**, 2191 (2005).
- ¹⁴B. T. Jonker, G. Kioseoglou, A. T. Hanbicki, C. H. Li, and P. E. Thompson, *Nat. Phys.* **3**, 542 (2007).
- ¹⁵S. P. Dash, S. Sharma, R. S. Patel, M. P. de Jong, and R. Jansen, *Nature* **462**, 491 (2009).
- ¹⁶B. T. Jonker, S. Erwin, A. Petrou, and A. G. Petukhov, *MRS Bull.* **28**, 740 (2003).
- ¹⁷R. A. de Groot, F. M. Mueller, P. G. van Engen, and K. H. J. Buschow, *Phys. Rev. Lett.* **50**, 2024 (1983).
- ¹⁸M. I. Katsnelson, V. Yu. Irkhin, L. Chioncel, A. I. Lichtenstein, and R. A. de Groot, *Rev. Mod. Phys.* **80**, 315 (2008).
- ¹⁹P. A. Dowben and R. Skomski, *J. Appl. Phys.* **95**, 7453 (2004).
- ²⁰Y. Sakuraba, M. Hattori, M. Oogane, Y. Ando, H. Kato, A. Sakuma, T. Miyazaki, and H. Kubota, *Appl. Phys. Lett.* **88**, 192508 (2006).
- ²¹T. Marukame, T. Ishikawa, K.-I. Matsuda, T. Uemura, and M. Yamamoto, *Appl. Phys. Lett.* **88**, 262503 (2006).
- ²²T. Ishikawa, T. Marukame, H. Kijima, K.-I. Matsuda, T. Uemura, M. Arita, and M. Yamamoto, *Appl. Phys. Lett.* **89**, 192505 (2006).
- ²³H. Sukegawa, W. Wang, R. Shan, T. Nakatani, K. Inomata, and K. Hono, *Phys. Rev. B* **79**, 184418 (2009).
- ²⁴T. M. Nakatani, T. Furubayashi, S. Kasai, H. Sukegawa, Y. K. Takahashi, S. Mitani, and K. Hono, *Appl. Phys. Lett.* **96**, 212501 (2010).
- ²⁵G. Bridoux, M. V. Costache, J. Van de Vondel, I. Neumann, and S. O. Valenzuela, *Appl. Phys. Lett.* **99**, 102107 (2011).
- ²⁶T. Kimura, N. Hashimoto, S. Yamada, M. Miyao, and K. Hamaya, *NPG Asia Mater.* **4**, e9 (2012).
- ²⁷K. Hamaya, N. Hashimoto, S. Oki, S. Yamada, M. Miyao, and T. Kimura, *Phys. Rev. B* **85**, 100404(R) (2012).
- ²⁸M. Ramsteiner, O. Brandt, T. Flissikowski, H. T. Grahn, M. Hashimoto, J. Herfort, and H. Kostial, *Phys. Rev. B* **78**, 121303(R) (2008).
- ²⁹S. Takahashi and S. Maekawa, *Phys. Rev. B* **67**, 052409 (2003).
- ³⁰B. L. Gyorffy, A. J. Pindor, J. Staunton, G. M. Stocks, and H. Winter, *J. Phys. F* **15**, 1337 (1985).
- ³¹R. Skomski and P. A. Dowben, *Europhys. Lett.* **58**, 544 (2002).
- ³²X. Waintal, E. B. Myers, P. W. Brouwer, and D. C. Ralph, *Phys. Rev. B* **62**, 12317 (2000).
- ³³P. A. Mello, P. Pereyra, and N. Kumar, *Ann. Phys. (NY)* **181**, 290 (1988).
- ³⁴C. W. J. Beenakker, *Rev. Mod. Phys.* **69**, 731 (1997).
- ³⁵P. W. Brouwer and C. W. J. Beenakker, *J. Math. Phys.* **37**, 4904 (1996).
- ³⁶In the notation of Ref. 32, matrices $R_{\lambda\mu}$ and $T_{\lambda\mu}$ correspond to the “hat matrices” generated by r'_1 and t_1 .
- ³⁷A. Brataas, Yu. V. Nazarov, and G. E. W. Bauer, *Phys. Rev. Lett.* **84**, 2481 (2000).
- ³⁸E. I. Rashba, *Eur. Phys. J. B* **29**, 513 (2002).
- ³⁹J. Bass and W. P. Pratt, *J. Phys.: Condens. Matter* **19**, 183201 (2007).
- ⁴⁰H. Y. T. Nguyen, R. Loloee, W. P. Pratt, and J. Bass, *Phys. Rev. B* **86**, 064413 (2012).
- ⁴¹S. Garzon, I. Žutić, and R. A. Webb, *Phys. Rev. Lett.* **94**, 176601 (2005).
- ⁴²L. Weber and E. Gmelin, *Appl. Phys. A* **53**, 136 (1991).
- ⁴³M. Ležaić, Ph. Mavropoulos, J. Enkovaara, G. Bihlmayer, and S. Blügel, *Phys. Rev. Lett.* **97**, 026404 (2006).
- ⁴⁴Ph. Mavropoulos, M. Ležaić, and S. Blügel, *Phys. Rev. B* **72**, 174428 (2005).
- ⁴⁵S. Chadov, T. Graf, K. Chadova, X. Dai, F. Casper, G. H. Fecher, and C. Felser, *Phys. Rev. Lett.* **107**, 047202 (2011).
- ⁴⁶T. N. Todorov, *Phys. Rev. B* **54**, 5801 (1996).