

Spintronics: Fundamentals and applications

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Spintronics, or spin electronics, involves the study of active control and manipulation of spin degrees of freedom in solid-state systems. This article reviews the current status of this subject, including both recent advances and well-established results. The primary focus is on the basic physical principles underlying the generation of carrier spin polarization, spin dynamics, and spin-polarized transport in semiconductors and metals. Spin transport differs from charge transport in that spin is a nonconserved quantity in solids due to spin-orbit and hyperfine coupling. The authors discuss in detail spin decoherence mechanisms in metals and semiconductors. Various theories of spin injection and spin-polarized transport are applied to hybrid structures relevant to spin-based devices and fundamental studies of materials properties. Experimental work is reviewed with the emphasis on projected applications, in which external electric and magnetic fields and illumination by light will be used to control spin and charge dynamics to create new functionalities not feasible or ineffective with conventional electronics.

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I. INTRODUCTION

A. Overview

Spintronics is a multidisciplinary field whose central theme is the active manipulation of spin degrees of free-

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dom in solid-state systems.¹ In this article the term spin stands for either the spin of a single electron \mathbf{s} , which can be detected by its magnetic moment $-g\mu_B \mathbf{s}$ (μ_B is the Bohr magneton and g is the electron g factor, in a solid generally different from the free electron value of $g_0 = 2.0023$), or the average spin of an ensemble of electrons, manifested by magnetization. The control of spin is then a control of either the population and the phase of the spin of an ensemble of particles, or a coherent spin manipulation of a single or a few-spin system. The goal of spintronics is to understand the interaction between the particle spin and its solid-state environments and to make useful devices using the acquired knowledge. Fundamental studies of spintronics include investigations of spin transport in electronic materials, as well as understanding spin dynamics and spin relaxation. Typical questions that are posed are (a) what is an effective way to polarize a spin system? (b) how long is the system able to remember its spin orientation? (c) how can spin be detected?

Generation of spin polarization usually means creating a nonequilibrium spin population. This can be achieved in several ways. While traditionally spin has been oriented using optical techniques in which circularly polarized photons transfer their angular momenta to electrons, for device applications electrical spin injection is more desirable. In electrical spin injection a magnetic electrode is connected to the sample. When the current drives spin-polarized electrons from the electrode to the sample, nonequilibrium spin accumulates there. The rate of spin accumulation depends on spin relaxation, the process of bringing the accumulated spin population back to equilibrium. There are several relevant mechanisms of spin relaxation, most involving spin-orbit coupling to provide the spin-dependent potential, in combination with momentum scattering providing a randomizing force. Typical time scales for spin relaxation in electronic systems are measured in nanoseconds, while the range is from pico to microseconds. Spin detection, also part of a generic spintronic scheme, typically relies on sensing the changes in the signals caused by the presence of nonequilibrium spin in the system. The common goal in many spintronic devices is to maximize the spin detection sensitivity to the point it detects not the spin itself, but changes in the spin states.

Let us illustrate the generic spintronic scheme on a prototypical device, the Datta-Das spin field effect transistor (SFET) (Datta and Das, 1990), depicted in Fig. 1. The scheme shows the structure of the usual FET, with a drain, a source, a narrow channel, and a gate for controlling the current. The gate either allows the current to flow (ON) or does not (OFF). The spin transistor is simi-

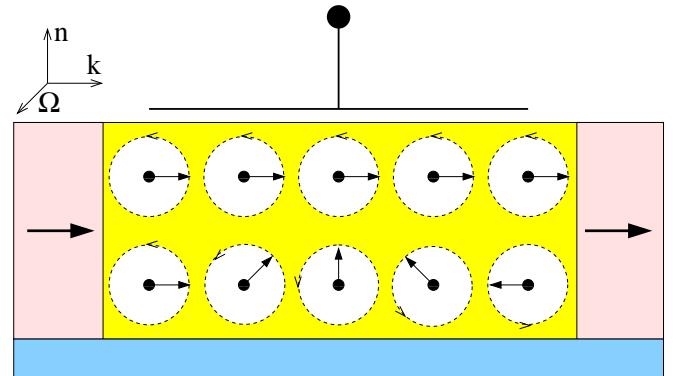


FIG. 1 Scheme of the Datta-Das spin field-effect transistor (SFET). The source (spin injector) and the drain (spin detector) are ferromagnetic metals or semiconductors, with parallel magnetic moments. The injected spin-polarized electrons with wave vector \mathbf{k} move ballistically along a quasi-one-dimensional channel formed by, for example, an In-GaAs/InAlAs heterojunction in a plane normal to \mathbf{n} . Electron spins precess about the precession vector $\boldsymbol{\Omega}$, which arises from spin-orbit coupling and which is defined by the structure and the materials properties of the channel. The magnitude of $\boldsymbol{\Omega}$ is tunable by the gate voltage V_G at the top of the channel. The current is large if the electron spin at the drain points in the initial direction (top row), for example, if the precession period is much larger than the time of flight, and small if the direction is reversed (bottom).

lar in that the result is also a control of the charge current through the narrow channel. The difference, however, is in the physical realization of the current control. In the Datta-Das SFET the source and the drain are ferromagnets acting as the injector and detector of the electron spin. The drain injects electrons with spins parallel to the transport direction. The electrons are transported ballistically through the channel. When they arrive at the drain, their spin is detected. In a simplified picture, the electron can enter the drain (ON) if its spin points in the same direction as the spin of the drain. Otherwise it is scattered away (OFF). The role of the gate is to generate an effective magnetic field (in the direction of $\boldsymbol{\Omega}$ in Fig. 1), arising from the spin-orbit coupling in the substrate material, from the confinement geometry of the transport channel, and the electrostatic potential of the gate. This effective magnetic field causes the electron spins to precess. By modifying the voltage, one can cause the precession to lead to either parallel or antiparallel (or anything between) electron spin at the drain, effectively controlling the current.

Even though the name *spintronics* is rather novel,² contemporary research in spintronics relies closely on a long tradition of results obtained in diverse areas of physics (for example, magnetism, semiconductor physics,

¹ While there are proposals for spintronic devices based on deoxyribonucleic acid (DNA) molecules (Zwolak and Di Ventra, 2002), the whole device, which includes electrodes, voltage/current source, etc., is still a solid-state system.

² The term was coined by S. A. Wolf in 1996, as a name for a DARPA initiative for novel magnetic materials and devices.

superconductivity, optics, and mesoscopic physics) and establishes new connections between its different sub-fields (Rashba, 2002d; Žutić, 2002). We review here both well-established results and the physical principles relevant to the present and future applications. Our strategy is to give a comprehensive view of what has been accomplished, focusing in detail on a few selected topics that we believe are representative for the broader subject within which they appear. For example, while discussing the generation of spin polarization, we survey many experimental and theoretical studies of both optical orientation and electrical spin injection and present a detailed and self-contained formalism of electrical spin injection. Similarly, when we discuss spin relaxation, we give a catalog of important work, while studying spin relaxation in the cases of Al and GaAs as representative of the whole field. Finally, in the section on spin devices we give detailed physical principles of several selected devices, such as, for example, the above-mentioned Datta-Das SFET.

There have been many other reviews written on spintronics, most focusing on a particular aspect of the field. We divide them here, for an easier orientation, into two groups, those that cover the emerging applications³ and those covering already well-established schemes and materials⁴. The latter group, often described as *magnetoelectronics* typically covers paramagnetic and ferromagnetic metals and insulators, which utilize magnetoresistive effects, realized, for example, as magnetic read heads in computer hard drives, nonvolatile magnetic random access memory (MRAM), and circuit isolators (Wang *et al.*, 2002). These more established aspects of spintronics have been also addressed in several books⁵ and will be discussed in another review,⁶ complementary to ours.

Spintronics also benefits from a large class of emerging materials, such as ferromagnetic semiconductors (Ohno, 1998; Pearton *et al.*, 2003), organic semiconductors (Dediu *et al.*, 2002), organic ferromagnets (Epstein, 2003; Pejaković *et al.*, 2002), high temperature superconductors (Goldman *et al.*, 1999), and carbon nanotubes (Tsukagoshi *et al.*, 1999; Zhao *et al.*, 2002), which can bring novel functionalities to the traditional devices. There is a continuing need for fundamental studies before the potential of spintronic applications is fully realized.

After an overview, Sec. I covers some basic histori-

cal and background material, part of which has already been extensively covered in the context of magnetoelectronics and will not be discussed further in this review. Techniques for generating spin polarization, focusing on optical spin orientation and electrical spin injection, are described in Sec. II. The underlying mechanisms responsible for the loss of spin orientation and coherence, which impose fundamental limits on the length and time scales in spintronic devices, are addressed in Sec. III. Spintronic applications and devices, with the emphasis on those based on semiconductors, are discussed in Sec. IV. The review concludes with a look at future prospects in Sec. V and with the table (Tab. II) listing the most common abbreviations used in the text.

B. History and background

1. Spin-polarized transport and magnetoresistive effects

In a pioneering work, Mott (1936a,b) provided a basis for our understanding of spin-polarized transport. Mott sought an explanation for an unusual behavior of resistance in ferromagnetic metals. He realized that at sufficiently low temperatures, where magnon scattering becomes vanishingly small, electrons of majority and minority spin, with magnetic moment parallel and antiparallel to the magnetization of a ferromagnet, respectively, do not mix in the scattering processes. The conductivity can then be expressed as the sum of two independent and unequal parts for two different spin projections—the current in ferromagnets is spin polarized. This is also known as the two-current model and has been extended by Campbell *et al.* (1967); Fert and Campbell (1968). It continues, in its modifications, to provide an explanation for various magnetoresistive phenomena (Valet and Fert, 1993).

Tunneling measurements played a key role in early experimental work on spin-polarized transport. Studying N/F/N junctions, where N was a nonmagnetic⁷ metal and F was an Eu-based ferromagnetic semiconductor (Kasuya and Yanase, 1968; Nagaev, 1983), revealed that I-V curves could be modified by an applied magnetic field (Esaki *et al.*, 1967) and show potential for developing a solid-state spin-filter. When unpolarized current is passed across a ferromagnetic semiconductor, the current becomes spin-polarized (Hao *et al.*, 1990; Moodera *et al.*, 1988).

A series of experiments (Tedrow and Meservey, 1971b, 1973, 1994) in ferromagnet/insulator/superconductor (F/I/S) junctions has unambiguously proved that the

³ Reviews on emerging application include those of (Das Sarma, 2001; Das Sarma *et al.*, 2000a,b, 2001, 2000c; Oestreich *et al.*, 2002; Rashba, 2002d; Wolf *et al.*, 2001; Wolf and Treger, 2000; Žutić, 2002; Žutić (Ed.), 2002).

⁴ Established schemes and materials are reviewed by (Ansermet, 1998; Bass and Pratt, Jr., 1999; Daughton *et al.*, 1999; Gijs and Bauer, 1997; Gregg *et al.*, 1997; Prinz, 1995, 1998; Stiles, 2004; Tedrow and Meservey, 1994).

⁵ See, for example, the books of (Chtchelkanova *et al.*, 2003; Hartman (Ed.), 2000; Hirota *et al.*, 2002; Levy and Mertig, 2002; Maekawa *et al.*, 2002; Parkin, 2002; Shinjo, 2002; Ziese and Thornton (Eds.), 2001)

⁶ In preparation by S. S. P. Parkin for Review Modern Physics.

⁷ Unless explicitly specified, we shall use the terms “nonmagnetic” and “paramagnetic” interchangeably, i.e., assume that they both refer to a material with no long-range ferromagnetic order and with Zeeman-split carrier spin subbands in an applied magnetic field.

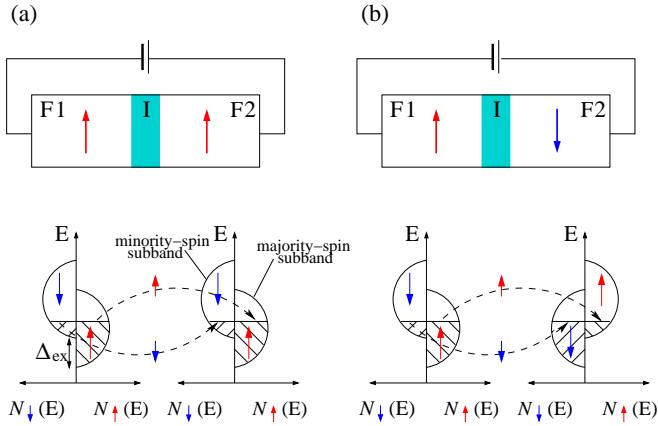


FIG. 2 Schematic illustration of electron tunneling in ferromagnet/insulator/ferromagnet (F/I/F) tunnel junctions: (a) Parallel and (b) antiparallel orientation of magnetizations with the corresponding spin-resolved density of the d states in ferromagnetic metals that have exchange spin splitting Δ_{ex} . Arrows in the two ferromagnetic regions are determined by the majority-spin subband. Dashed lines depict spin-conserved tunneling.

tunneling current remains spin-polarized even outside of the ferromagnetic region.⁸ The Zeeman-split quasiparticle density of states in a superconductor (Fulde, 1973; Tedrow *et al.*, 1970) was used as a detector of spin polarization of conduction electrons in various magnetic materials. Jullière (1975) measured tunneling conductance of F/I/F junctions, where I was an amorphous Ge. By adopting the Tedrow and Meservey (1971b, 1973) analysis of the tunneling conductance from F/I/S to the F/I/F junctions, Jullière (1975) formulated a model for a change of conductance between the parallel ($\uparrow\uparrow$) and antiparallel ($\uparrow\downarrow$) magnetization in the two ferromagnetic regions F1 and F2, as depicted in Fig. 2. The corresponding tunneling magnetoresistance⁹ (TMR) in an F/I/F magnetic tunnel junction (MTJ) is defined as

$$TMR = \frac{\Delta R}{R_{\uparrow\uparrow}} = \frac{R_{\uparrow\downarrow} - R_{\uparrow\uparrow}}{R_{\uparrow\uparrow}} = \frac{G_{\uparrow\downarrow} - G_{\uparrow\uparrow}}{G_{\uparrow\uparrow}}, \quad (1)$$

where conductance G and resistance $R=1/G$ are labeled by the relative orientations of the magnetizations in F1 and F2 (it is possible to change the relative orientations, between $\uparrow\uparrow$ and $\uparrow\downarrow$, even at small applied magnetic fields ~ 10 G). TMR is a particular manifestation of a magnetoresistance (MR) that yields a change of electrical resistance in the presence of an external magnetic field.¹⁰ Historically, the anisotropic MR in bulk ferromagnets such

as Fe and Ni was discovered first, dating back to the experiments of Lord Kelvin (Thomson, 1857). Due to spin-orbit interaction, electrical resistivity changes with the relative direction of the charge current (for example, parallel or perpendicular) with respect to the direction of magnetization.

Within Jullière's model, which assumes constant tunneling matrix elements and that electrons tunnel without spin flip, Eq. (1) yields

$$TMR = \frac{2P_1P_2}{1 - P_1P_2}, \quad (2)$$

where the polarization $P_i = (\mathcal{N}_{Mi} - \mathcal{N}_{mi})/(\mathcal{N}_{Mi} + \mathcal{N}_{mi})$ is expressed in terms of the spin-resolved density of states \mathcal{N}_{Mi} and \mathcal{N}_{mi} , for majority and minority spin in F_i , respectively. Conductance in Eq. (1) can then be expressed as (Maekawa and Gäfvert, 1982) $G_{\uparrow\uparrow} \sim \mathcal{N}_{M1}\mathcal{N}_{M2} + \mathcal{N}_{m1}\mathcal{N}_{m2}$ and $G_{\uparrow\downarrow} \sim \mathcal{N}_{M1}\mathcal{N}_{m2} + \mathcal{N}_{m1}\mathcal{N}_{M2}$ to give Eq. (2).¹¹ While the early results of Jullière (1975) were not confirmed, TMR at 4.2 K was observed using NiO as a tunnel barrier by Maekawa and Gäfvert (1982).

The prediction of Jullière's model illustrates the spin-valve effect: the resistance of a device can be changed by manipulating the relative orientation of the magnetizations \mathbf{M}_1 and \mathbf{M}_2 , in F1 and F2, respectively. Such orientation can be preserved even in the absence of a power supply and the spin-valve effect,¹² later discovered in multilayer structures displaying the giant magnetoresistance (GMR) effect,¹³ (Baibich *et al.*, 1988; Binnsch *et al.*, 1989) can be used for nonvolatile memory applications (Hartman (Ed.), 2000; Hirota *et al.*, 2002; Parkin, 2002). GMR structures are often classified according whether the current flows parallel (CIP) or perpendicular (CPP) to the interfaces between the different layers, as depicted in Fig. 3. Most of the GMR applications use the CIP geometry, while the CPP version, first realized by (Pratt, Jr. *et al.*, 1991), is easier to analyze theoretically (Gijs and Bauer, 1997; Levy and Mertig, 2002) and relates to the physics of the TMR effect (Mathon and Umerski, 1997). The size of magnetoresistance in the GMR structures can be expressed analogously to Eq. (1), where parallel and antiparallel orientations of the magnetizations in the two ferromagnetic regions are often denoted by "P" and "AP," respectively (instead of $\uparrow\uparrow$ and $\uparrow\downarrow$). Realization of a large room temperature GMR (Parkin *et al.*, 1991a,b) enabled a quick

⁸ It has been shown that electrons photoemitted from ferromagnetic gadolinium remain spin polarized (Busch *et al.*, 1969).

⁹ Starting with Jullière (1975) an equivalent expression $(G_{\uparrow\uparrow} - G_{\uparrow\downarrow})/G_{\uparrow\uparrow}$ has also been used by different authors and often referred to as junction magnetoresistance (JMR) (Moodera and Mathon, 1999).

¹⁰ The concept of TMR was proposed independently by R. C.

Barker in 1975 [see Meservey *et al.* (1983)] and by Slonczewski (1976), who envisioned its use for magnetic bubble memory (Parkin, 2002).

¹¹ In IV we address some limitations of the Jullière's model and its potential ambiguities to identify precisely which spin polarization is actually measured.

¹² The term was coined by Dieny *et al.* (1991) in the context of GMR, by invoking an analogy with the physics of the TMR.

¹³ The term "giant" reflected the magnitude of the effect (more than $\sim 10\%$), as compared to the better known anisotropic magnetoresistance ($\sim 1\%$).

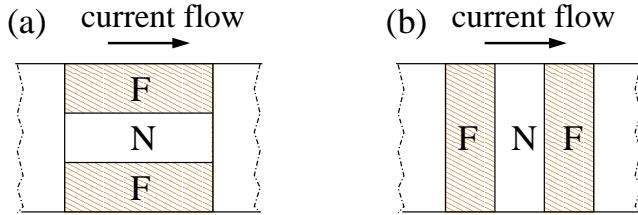


FIG. 3 Schematic illustration of (a) the current in plane (CIP) (b) the current perpendicular to the plane (CPP) giant magnetoresistance geometry.

transition from basic physics to commercial applications in magnetic recording (Parkin *et al.*, 2003).

One of the keys to the success of the MR-based applications is their ability to control¹⁴ the relative orientation of \mathbf{M}_1 and \mathbf{M}_2 . An interesting realization of such control was proposed independently by Berger (1996) and Slonczewski (1996). While in GMR or TMR structures the relative orientation of magnetizations will affect the flow of spin-polarized current, they predicted a reverse effect. The flow of spin-polarized current can transfer angular momentum from carriers to ferromagnet and alter the orientation of the corresponding magnetization, even in the absence of an applied magnetic field. This phenomenon, known as spin-transfer torque, has since been extensively studied both theoretically and experimentally (Bazaliy *et al.*, 1998; Myers *et al.*, 1999; Stiles and Zangwill, 2002; Sun, 2000; Tsoi *et al.*, 1998; Waintal *et al.*, 2000) and current-induced magnetization reversal was demonstrated at room temperature (Katine *et al.*, 2000). It was also shown that the magnetic field generated by passing the current through a CPP GMR device could produce room temperature magnetization reversal (Bussmann *et al.*, 1999). In the context of ferromagnetic semiconductors additional control of magnetization was demonstrated optically (by shining light) (Boukari *et al.*, 2002; Koshihara *et al.*, 1997; Oiwa *et al.*, 2002) and electrically (by applying gate voltage) (Boukari *et al.*, 2002; Ohno *et al.*, 2000a; Park *et al.*, 2002) to perform switching between the ferromagnetic and paramagnetic states.

Jullière's model also justifies the continued quest for highly spin-polarized materials – they would provide large magnetoresistive effects, desirable for device applications. In an extreme case, spins would be completely polarized even in the absence of magnetic field. Numerical support for the existence of such materials—the so called half-metallic ferromagnets¹⁵ was provided by de Groot *et al.* (1983b), and these materials were reviewed by Pickett and Moodera (2001). In addition to ferromagnets, such as CrO_2 (Parker *et al.*, 2002; Soulen Jr. *et al.*,

1998) and manganite perovskites (Park *et al.*, 1998a), there is evidence for high spin polarization in III-V ferromagnetic semiconductors like $(\text{Ga},\text{Mn})\text{As}$ (Braden *et al.*, 2003; Panguluri *et al.*, 2003a). The challenge remains to preserve such spin polarization above room temperature and in junctions with other materials, since the surface (interface) and bulk magnetic properties can be significantly different (Falicov *et al.*, 1990; Fisher, 1967; Mills, 1971).

While many existing spintronic applications (Hartman (Ed.), 2000; Hirota *et al.*, 2002) are based on the GMR effects, the discovery of large room-temperature TMR (Miyazaki and Tezuka, 1995; Moodera *et al.*, 1995) has renewed interest in the study of magnetic tunnel junctions which are now the basis for the several MRAM prototypes¹⁶ (Parkin *et al.*, 1999; Tehrani *et al.*, 2000). Future generations of magnetic read heads are expected to use MTJ's instead of CIP GMR. To improve the switching performance of related devices it is important to reduce the junction resistance, which determines the RC time constant of the MTJ cell. Consequently, semiconductors, which would provide a lower tunneling barrier than the usually employed oxides, are being investigated both as the non-ferromagnetic region in MTJ's and as the basis for an all-semiconductor junction that would demonstrate large TMR at low temperatures (Tanaka, 2002; Tanaka and Higo, 2001). Another desirable property of semiconductors has been demonstrated by the extraordinary large room-temperature MR in hybrid structures with metals reaching 750 000% at a magnetic field of 4 T (Solin *et al.*, 2000) which could lead to improved magnetic read heads (Moussa *et al.*, 2003; Solin *et al.*, 2002). MR effects of similar magnitude have also been found in hybrid metal/semiconductor granular films (Akainaga, 2002). Another approach to obtaining large room-temperature magnetoresistance (> 100% at $B \sim 100$ G) is to fabricate ferromagnetic regions separated by a nano-size contact. For simplicity, such a structure could be thought of as the limiting case of the CPP GMR scheme in Fig. 3(b). This behavior, also known as ballistic magnetoresistance, has already been studied in a large number of materials and geometries (Bruno, 1999; Chung *et al.*, 2002; Garcia *et al.*, 1999; Imamura *et al.*, 2000; Tatara *et al.*, 1999; Versluijs *et al.*, 2001).

2. Spin injection and optical orientation

Many materials in their ferromagnetic state can have a substantial degree of *equilibrium* carrier spin polarization. However, as illustrated in Fig. 1, this alone is usually not sufficient for spintronic applications, which typically require current flow and/or manipulation of the *nonequilibrium*

¹⁴ For example, with small magnetic field (Parkin, 2002) or at high switching speeds (Schumacher *et al.*, 2003a,b).

¹⁵ Near the Fermi level they behave as metals only for one spin, the density of states vanishes completely for the other spin.

¹⁶ Realization of the early MRAM proposals used the effect of anisotropic magnetoresistance (Pohm *et al.*, 1987, 1988).

rium spin (polarization).¹⁷ The importance of generating *nonequilibrium* spin is not limited to device applications; it can also be used as a sensitive spectroscopic tool to study a wide variety of fundamental properties ranging from spin-orbit and hyperfine interactions (Meier and Zakharchenya (Eds.), 1984) to the pairing symmetry of high temperature superconductors (Ngai *et al.*, 2004; Tsuei and Kirtley, 2000; Vas'ko *et al.*, 1997; Wei *et al.*, 1999) and the creation of spin-polarized beams to measure parity violation in high energy physics (Pierce and Celotta, 1984).

Nonequilibrium spin is the result of some source of pumping arising from transport, optical, or resonance methods. Once the pumping is turned off the spin will return to its equilibrium value. While for most applications it is desirable to have long spin relaxation times, it has been demonstrated that short spin relaxation times are useful in the implementation of fast switching (Nishikawa *et al.*, 1995).

Electrical spin injection, an example of a transport method for generating nonequilibrium spin, has already been realized experimentally by Clark and Feher (1963), who drove a direct current through a sample of InSb in the presence of constant applied magnetic field. The principle was based on the Feher effect,¹⁸ in which the hyperfine coupling between the electron and nuclear spins, together with different temperatures representing electron velocity and electron spin populations, is responsible for the dynamical nuclear polarization (Slichter, 1989).¹⁹ Motivated by the work of Clark and Feher (1963), Tedrow and Meservey (1971b, 1973), and the principle of optical orientation (Meier and Zakharchenya (Eds.), 1984), Aronov (1976a,b) and Aronov and Pikus (1976) established several key concepts in electrical spin injection

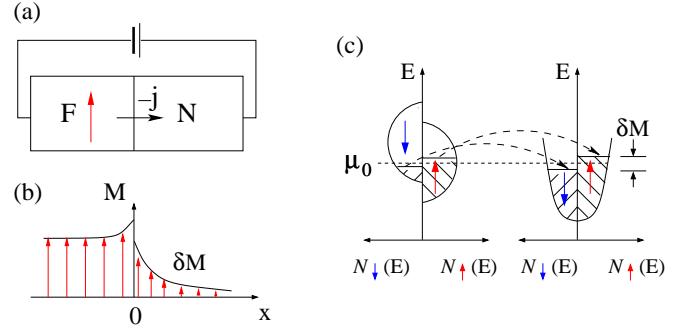


FIG. 4 Pedagogical illustration of the concept of electrical spin injection from a ferromagnet (F) into a normal metal (N). Electrons flow from F to N: (a) schematic device geometry; (b) magnetization M as a function of position. Nonequilibrium magnetization δM (spin accumulation) is injected into a normal metal; (c) contribution of different spin-resolved densities of states to charge and spin transport across the F/N interface. Unequal filled levels in the density of states depict spin-resolved electrochemical potentials different from the equilibrium value μ_0 .

from ferromagnets into metals, semiconductors²⁰ and superconductors. When a charge current flowed across the F/N junction (Fig. 4) Aronov (1976b) predicted that spin-polarized carriers in a ferromagnet would contribute to the net current of magnetization entering the nonmagnetic region and would lead to nonequilibrium magnetization δM , depicted in Fig. 4(b), with the spatial extent given by the spin diffusion length (Aronov, 1976b; Aronov and Pikus, 1976).²¹ Such δM , which is also equivalent to a *nonequilibrium* spin accumulation, was first measured in metals by Johnson and Silsbee (1985, 1988d). In the steady state δM is realized as the balance between spins added by the magnetization current and spins removed by spin relaxation.²²

Generation of nonequilibrium spin polarization and spin accumulation is also possible by optical methods known as optical orientation or optical pumping. In optical orientation, the angular momentum of absorbed circularly polarized light is transferred to the medium. Electron orbital momenta are directly oriented by light

¹⁷ Important exceptions are tunneling devices operating at low bias and near *equilibrium* spin. Equilibrium polarization and the current flow can be potentially realized, for example, in spin-triplet superconductors and thin-film ferromagnets (König *et al.*, 2001), accompanied by dissipationless spin currents. Using an analogy with the quantum Hall effect, it has been suggested that the spin-orbit interaction could lead to dissipationless spin currents in hole-doped semiconductors (Murakami *et al.*, 2003). Rashba (2003b) has pointed out that similar dissipationless spin currents in thermodynamic equilibrium, due to spin-orbit interaction, are not transport currents which could be employed for transporting spins and spin injection. It is also instructive to compare several earlier proposals that use spin-orbit coupling to generate spin currents, discussed in Sec. II.A.

¹⁸ The importance and possible applications of the Feher effect (Feher, 1959) to polarize electrons was discussed by (Das Sarma *et al.*, 2000b; Suhl, 2002).

¹⁹ Such an effect can be thought of as a generalization of the Overhauser effect (Overhauser, 1953b) in which the use of a resonant microwave excitation causes the spin relaxation of the nonequilibrium electron population through hyperfine coupling to lead to the spin polarization of nuclei. Feher (1959) suggested several other methods, instead of microwave excitation, that could produce a nonequilibrium electron population and yield a dynamical polarization of nuclei [see also Weger (1963)].

²⁰ In an earlier work spin injection of minority carriers was proposed in a ferromagnet/insulator/p-type semiconductor structure. Measuring polarization of electroluminescence was suggested as a technique for detecting injection of polarized carriers in a semiconductor (Scifres *et al.*, 1973).

²¹ Supporting the findings of Clark and Feher (1963), Aronov calculated that the electrical spin injection would polarize nuclei and lead to a measurable effect in the electron spin resonance (ESR). Several decades later related experiments on spin injection are also examining other implications of dynamical nuclear polarization (Johnson, 2000; Strand *et al.*, 2003).

²² The spin diffusion length is an important quantity for CPP GMR. The thickness of the N region in Fig. 3 should not exceed the spin diffusion length, otherwise the information on the orientation of the magnetization in F1 will not be transferred to the F2 region.

and through spin-orbit interaction electron spins become polarized. In II.B we focus on the optical orientation in semiconductors, a well-established technique (Meier and Zakharchenya (Eds.), 1984). In a pioneering work Lampel (1968) demonstrated that spins in silicon can be optically oriented (polarized). This technique is derived from optical pumping proposed by Kastler (1950) in which optical irradiation changes the relative populations within the Zeeman and the hyperfine levels of the ground states of atoms. While there are similarities with previous studies of free atoms (Cohen-Tannoudji and Kostler, 1966; Happer, 1972), optical orientation in semiconductors has important differences related to the strong coupling between the electron and nuclear spin and macroscopic number of particles (Hermann *et al.*, 1985; Meier and Zakharchenya (Eds.), 1984; Paget *et al.*, 1977). Polarized nuclei can exert large magnetic fields (~ 5 T) on electrons. In bulk III-V semiconductors, such as GaAs, optical orientation can lead to 50% polarization of electron density which could be further enhanced in quantum structures of reduced dimensionality or by applying a strain. A simple reversal in the polarization of the illuminating light (from positive to negative helicity) also reverses the sign of the electron density polarization. Combining these properties of optical orientation with the semiconductors tailored to have a negative electron affinity allows photoemission of spin-polarized electrons to be used as a powerful detection technique in high-energy physics and for investigating surface magnetism (Pierce and Celotta, 1984).

II. GENERATION OF SPIN POLARIZATION

A. Introduction

Transport, optical, and resonance methods (as well as their combination) have all been used to create nonequilibrium spin. After introducing the concept of spin polarization in solid-state systems we give a pedagogical picture of electrical spin injection and detection of polarized carriers. While electrical spin injection and optical orientation will be discussed in more detail later in this section, we also survey here several other techniques for polarizing carriers.

Spin polarization not only of electrons, but also of holes, nuclei, and excitations can be defined as

$$P_X = X_s/X, \quad (3)$$

the ratio of the difference $X_s = X_\lambda - X_{-\lambda}$ and the sum $X = X_\lambda + X_{-\lambda}$, of the spin-resolved λ components for a particular quantity X . To avoid ambiguity as to what precisely is meant by spin polarization both the choice of the spin-resolved components and the relevant physical quantity X need to be specified. Conventionally, λ is taken to be \uparrow or $+$ (numerical value +1) for spin up, \downarrow or $-$ (numerical value -1) for spin down, with respect

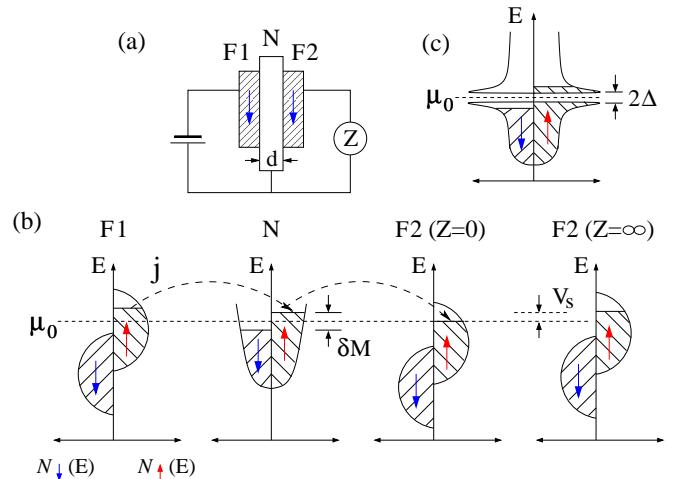


FIG. 5 Spin injection, spin accumulation, and spin detection: (a) two idealized completely polarized ferromagnets F1 and F2 (the spin down density of states \mathcal{N}_\downarrow is zero at the energy of electrochemical potential $E = \mu_0$) with parallel magnetizations are separated by the nonmagnetic region N; (b) density-of-states diagrams for spin injection from F1 into N, accompanied by the spin accumulation-generation of the nonequilibrium magnetization δM . At F2 in the limit of low impedance ($Z=0$) spin is detected by measuring the spin-polarized current across the N/F2 interface. In the limit of high impedance ($Z = \infty$) spin is detected by measuring the voltage $V_s \sim \delta M$ developed across the N/F2 interface; (c) spin accumulation in a device in which a superconductor (with the superconducting gap Δ) is occupying the region between F1 and F2.

to the chosen axis of quantization.²³ In ferromagnetic metals it is customary to refer to \uparrow (\downarrow) as carriers with magnetic moment parallel (antiparallel) to the magnetization or, equivalently, as carriers with majority (minority) spin (Tedrow and Meservey, 1973). In semiconductors the terms majority and minority usually refer to relative populations of the carriers while \uparrow or $+$ and \downarrow or $-$ correspond to the quantum numbers m_j with respect to the z -axis taken along the direction of the light propagation or along the applied magnetic field (Jonker *et al.*, 2003b; Meier and Zakharchenya (Eds.), 1984). It is important to emphasize that both the magnitude and the sign of the spin polarization in Eq. (3) depends of the choice of X , relevant to the detection technique employed, say optical vs. transport and bulk vs. surface measurements (Jonker *et al.*, 2003b; Mazin, 1999). Even in the same homogeneous material the measured P_X can vary for different X , and it is crucial to identify which physical quantity—charge current, carrier density, conductivity, or the density of states—is being measured experimentally.

The spin polarization of electrical current or carrier

²³ For example, along the spin angular momentum, applied magnetic field, magnetization, or direction of light propagation.

density, generated in a nonmagnetic region, is typically used to describe the efficiency of electrical spin injection. Silsbee (1980) suggested that the nonequilibrium density polarization in the N region, or equivalently the nonequilibrium magnetization, acts as the source of spin electromotive force (EMF) and produces a measurable “spin-coupled” voltage $V_s \propto \delta M$. Using this concept, also referred to as *spin-charge coupling*, Silsbee (1980) proposed a detection technique consisting of two ferromagnets F1 and F2 (see Fig. 5) separated by a nonmagnetic region.²⁴ F1 serves as the spin injector (spin aligner) and F2 as the spin detector. This could be called the polarizer-analyzer method, the optical counterpart of the transmission of light through two optical linear polarizers. From Fig. 5 it follows that the reversal of the magnetization direction in one of the ferromagnets would lead either to $V_s \rightarrow -V_s$, in an open circuit (in the limit of large impedance Z), or to the reversal of charge current $j \rightarrow -j$, in a short circuit (at small Z), a consequence of Silsbee-Johnson spin-charge coupling (Johnson and Silsbee, 1987, 1988a; Silsbee, 1980). Correspondingly, as discussed in the following sections, the spin injection could be detected through the spin accumulation signal either as a voltage or the resistance change when the magnetizations in F1 and F2 are changed from parallel to antiparallel alignment.

Since the experiments demonstrating the spin accumulation of conduction electrons in metals (Johnson and Silsbee, 1985), spin injection has been realized in a wide range of materials. While in Sec. II.C we focus on related theoretical work motivated by potential applications, experiments on spin injection have also stimulated proposals for examining the fundamental properties of electronic systems.²⁵

The generation of nonequilibrium spin polarization has a long tradition in magnetic resonance methods (Abragam, 1961; Slichter, 1989). However, transport methods to generate carrier spin polarization are not limited to electrical spin injection. For example, they also include scattering of unpolarized electrons in the presence of spin-orbit coupling (Kessler, 1976; Mott and Massey, 1965) and in materials that lack the inversion symmetry (Levitov *et al.*, 1984), adiabatic (Mucciolo *et al.*, 2002; Sharma and Chamon, 2003; Watson *et al.*, 2003) and nonadiabatic quantum spin pumping (Zheng *et al.*, 2003) [for an instructive description of parametric pumping see (Brouwer, 1998)], and the proximity effects (Ciuti *et al.*, 2002a).

²⁴ A similar geometry was also proposed independently by de Groot *et al.* (1983a), where F1 and F2 were two half-metallic ferromagnets with the goal of implementing spin-based devices to amplify and/or switch current.

²⁵ For example, studies probing the spin-charge separation in the non-Fermi liquids have been proposed by (Balents and Egger, 2000, 2001; Kivelson and Rokhsar, 1990; Si, 1997, 1998; Zhao and Hershfield, 1995). Spin and charge are carried by separate excitations and can lead to spatially separated spin and charge currents (Kivelson and Rokhsar, 1990).

It would be interesting to know what the limits are on the magnitude of various spin polarizations. Could we have a completely polarized current [$P_j \rightarrow \infty$, see Eq. (3)], with only a spin current ($j_\uparrow - j_\downarrow$) and no charge current ($j_\uparrow + j_\downarrow = 0$)? While it is tempting to recall the Stern-Gerlach experiment and try to set up magnetic drift through inhomogeneous magnets (Kessler, 1976), this would most likely work only as a transient effect (Fabian and Das Sarma, 2002). It was proposed already by D'yakonov and Perel' (1971a,c) that a transverse spin current (and transverse spin polarization in a closed sample) would form as a result of spin-orbit coupling-induced skew scattering in the presence of a longitudinal electric field. This interesting effect, also called the *spin Hall effect* (Hirsch, 1999; Zhang, 2000), has yet to be demonstrated. An alternative scheme for producing pure spin currents was proposed by Bhat and Sipe (2000), motivated by the experimental demonstration of phase coherent control of charge currents (Atanasov *et al.*, 1996; Haché *et al.*, 1997) and carrier population (Fraser *et al.*, 1999). A quantum-mechanical interference between one- and two-photon absorptions of orthogonal linear polarizations creates an opposite ballistic flow of spin up and spin down electrons in a semiconductor. Only a spin current can flow, without a charge current, as demonstrated by Stevens *et al.* (2003) and Hübner *et al.* (2003), who were able to achieve coherent control of the spin current direction and magnitude by the polarization and the relative phase of two exciting laser light fields.

Charge current also can be driven by circularly polarized light (Ivchenko and Pikus, 1997). Using the principles of optical orientation (see Sec. I.B.2 and further discussion in Sec. II.B) in semiconductors of reduced dimensionality or lower symmetry, both the direction and the magnitude of a generated charge current can be controlled by circular polarization of the light. This is called the circular photo-voltaic effect (Ganichev and Prettl, 2003), which can be viewed as a transfer of the angular momentum of photons to directed motion of electrons. This could also be called a spin corkscrew effect, since a nice mechanical analog is a corkscrew whose rotation generates linear directed motion. A related effect, in which the photocurrent is driven, is called the spin-galvanic effect (Ganichev and Prettl, 2003). The current here is caused by the difference in spin-flip scattering rates for electrons with different spin states in some systems with broken inversion symmetry. A comprehensive survey of the related effects from the circular photo-galvanic effect (Asnin *et al.*, 1979) to recent demonstrations in semiconductor quantum wells (Ganichev *et al.*, 2002a,b, 2001, 2003)] is given by Ganichev and Prettl (2003).

There is a wide range of recent theoretical proposals for devices that would give rise to a spin electromotive force (Brataas *et al.*, 2002; Governale *et al.*, 2003; Long *et al.*, 2003; Mal'shukov *et al.*, 2003; Ting and Cartoixà, 2003; Žutić *et al.*, 2001a,b), often referred to as spin(-polarized) pumps, cells, or batteries. However, even when it is feasible to generate pure spin current, this does not directly

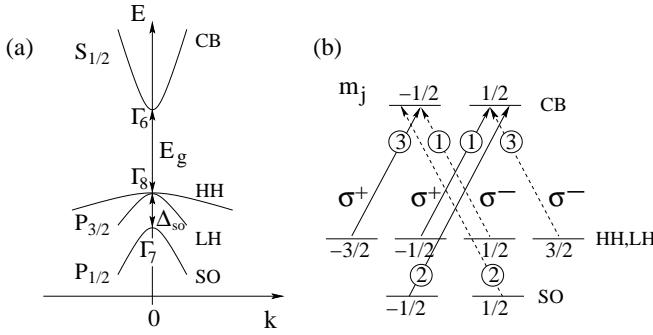


FIG. 6 (a) Interband transitions in GaAs: (a) schematic band structure of GaAs near the center of the Brillouin zone (Γ point), where E_g is the band gap and Δ_{so} is the spin-orbit splitting; CB, conduction band; HH, valence heavy hole; LH, light hole; SO, spin-orbit-split-off subbands; $\Gamma_{6,7,8}$ are the corresponding symmetries at the $k = 0$ point, more precisely, the irreducible representations of the tetrahedron group T_d (Ivchenko and Pikus, 1997); (b) selection rules for interband transitions between m_j sublevels for circularly polarized light σ^+ and σ^- (positive and negative helicity). The circled numbers denote the relative transition intensities that apply for both excitations (depicted by the arrows) and radiative recombinations.

imply that it would be dissipationless. In the context of superconductors, it has been shown that Joule heating can arise from pure spin current flowing through a Josephson junction (Takahashi *et al.*, 2001).

B. Optical spin orientation

In a semiconductor the photoexcited spin-polarized electrons and holes exist for the time τ before they recombine. If a fraction of the carriers' initial orientation survives longer than the recombination time, that is, if $\tau < \tau_s$,²⁶ where τ_s is the spin relaxation time (see Sec. III), the luminescence (recombination radiation) will be partially polarized. By measuring the circular polarization of the luminescence it is possible to study the spin dynamics of the nonequilibrium carriers in semiconductors (Oestreich *et al.*, 2002) and to extract such useful quantities as the spin orientation, the recombination time, or the spin relaxation time of the carriers (Ekimov and Safarov, 1970; Garbuzov *et al.*, 1971; Meier and Zakharchenya (Eds.), 1984; Parsons, 1969).

We illustrate the basic principles of optical orientation by the example of GaAs which is representative of a large class of III-V and II-VI zincblende semiconductors. The band structure is depicted in Fig. 6(a). The band gap is $E_g = 1.52$ eV at $T = 0$ K, while the spin split-off

²⁶ In Si this condition is not fulfilled. Instead of measuring the luminescence polarization, Lampel (1968) has used NMR to detect optical spin orientation.

Symmetry $ J, m_j\rangle$	Wave function
Γ_6	$ 1/2, 1/2\rangle S\uparrow\rangle$
	$ 1/2, -1/2\rangle S\downarrow\rangle$
Γ_7	$ 1/2, 1/2\rangle -(1/3)^{1/2}[(X+iY)\downarrow -Z\uparrow]\rangle$
	$ 1/2, -1/2\rangle (1/3)^{1/2}[(X-iY)\uparrow +Z\downarrow]\rangle$
Γ_8	$ 3/2, 3/2\rangle (1/2)^{1/2}(X+iY)\uparrow\rangle$
	$ 3/2, 1/2\rangle (1/6)^{1/2}[(X+iY)\downarrow +2Z\uparrow]\rangle$
	$ 3/2, -1/2\rangle -(1/6)^{1/2}[(X-iY)\uparrow -2Z\downarrow]\rangle$
	$ 3/2, -3/2\rangle (1/2)^{1/2}(X-iY)\downarrow\rangle$

TABLE I Angular and spin part of the wave function at Γ .

band is separated from the light and heavy hole bands by $\Delta_{so} = 0.34$ eV. We denote the Bloch states according to the total angular momentum J and its projection onto the positive z axis m_j : $|J, m_j\rangle$. Expressing the wave functions with the symmetry of s , p_x , p_y , and p_z orbitals as $|S\rangle$, $|X\rangle$, $|Y\rangle$, and $|Z\rangle$, respectively, the band wave functions can be written as listed in Table I (Pierce and Meier, 1976) [with minor typos removed, see also (Kittel, 1963)].

To obtain the excitation (or recombination) probabilities consider photons arriving in the z direction. Let σ^\pm represent the helicity of the exciting light. When we represent the dipole operator corresponding to the σ^\pm optical transitions as²⁷ $\propto (X \pm iY) \propto Y_l^{\pm 1}$, where Y_l^m is the spherical harmonic, it follows from Table I that

$$\frac{|\langle 1/2, -1/2|Y_1^1|3/2, -3/2\rangle|^2}{|\langle 1/2, 1/2|Y_1^1|3/2, -1/2\rangle|^2} = 3 \quad (4)$$

for the relative intensity of the σ^+ transition between the heavy ($|m_j = 3/2|$) and the light ($|m_j = 1/2|$) hole subbands and the conduction band. Other transitions are analogous. The relative transition rates are indicated in Fig. 6(b). The same selection rules apply to the optical orientation of shallow impurities (Ekimov and Safarov, 1970; Parsons, 1969).

The spin polarization of the excited electrons²⁸ depends on the photon energy $\hbar\omega$. For $\hbar\omega$ between E_g and $E_g + \Delta_{so}$, only the light and heavy hole subbands contribute. Denoting by n_+ and n_- the density of electrons polarized parallel ($m_j = 1/2$) and antiparallel ($m_j = -1/2$) to the direction of light propagation, we

²⁷ For an outgoing light in the $-z$ direction the helicities are reversed.

²⁸ Although holes are initially polarized too, they lose spin orientation very fast, on the time scale of the momentum relaxation time (see Sec. III.D.1). However, it was suggested that manipulating hole spin by short electric field pulses, between momentum scattering events, could be useful for ultrafast spintronics applications (Dargys, 2002).

define the spin polarization as (see Sec. II.A)

$$P_n = (n_+ - n_-)/(n_+ + n_-). \quad (5)$$

For our example of the zincblende structure,

$$P_n = (1 - 3)/(3 + 1) = -1/2 \quad (6)$$

is the spin polarization at the moment of photoexcitation. The spin is oriented against the direction of light propagation, since there are more transitions from the heavy hole than from the light hole subbands. The circular polarization of the luminescence is defined as

$$P_{\text{circ}} = (I^+ - I^-)/(I^+ + I^-), \quad (7)$$

where I^\pm is the radiation intensity for the helicity σ^\pm . The polarization of the σ^+ photoluminescence is then

$$P_{\text{circ}} = \frac{(n_+ + 3n_-) - (3n_+ + n_-)}{(n_+ + 3n_-) + (3n_+ + n_-)} = -\frac{P_n}{2} = \frac{1}{4}. \quad (8)$$

If the excitation involves transitions from the spin split-off band, that is, if $\hbar\omega \gg E_g + \Delta_{so}$, the electrons will not be spin polarized ($P_n = P_{\text{circ}} = 0$), underlining the vital role of spin-orbit coupling for spin orientation. On the other hand, Fig. 6 suggests that a removal of the heavy/light hole degeneracy can substantially increase P_n (D'yakonov and Perel', 1984a), up to the limit of complete spin polarization. An increase in P_n and P_{circ} in GaAs strained due to a lattice mismatch with a substrate, or due to confinement in quantum well heterostructures, has indeed been demonstrated (Oskotskij *et al.*, 1997; Vasilev *et al.*, 1993), detecting P_n greater than 0.9.

While photoexcitation with circularly polarized light creates spin-polarized electrons, the nonequilibrium spin decays due to both carrier recombination and spin relaxation. The steady-state degree of spin polarization depends on the balance between the spin excitation and decay. Sometimes a distinction is made (Meier and Zakharchenya (Eds.), 1984; Pierce and Meier, 1976) between the terms optical *spin orientation* and optical *spin pumping*. The former term is used in relation to the minority carriers (such as electrons in p-doped samples) and represents the orientation of the excited carriers. The latter term is reserved for the majority carriers (electrons in n-doped samples), representing spin polarization of the “ground” state. Both spin orientation and spin pumping were demonstrated in the early investigations on p-GaSb (Parsons, 1969) and p- and n-Ga_{0.7}Al_{0.3}As (Ekimov and Safarov, 1970, 1971; Zakharchenya *et al.*, 1971). Unless specified otherwise, we shall use the term optical orientation to describe both spin orientation and spin pumping.

To derive the steady-state expressions for the spin polarization due to optical orientation, consider the simple model of carrier recombination and spin relaxation (see Sec. IV.A.4) in a homogeneously doped semiconductor. The balance between direct electron-hole recombination and optical pair creation can be written as

$$r(np - n_0 p_0) = G, \quad (9)$$

where r measures the recombination rate, the electron and hole densities are n and p , with index zero denoting the equilibrium values, and G is the electron-hole photoexcitation rate. Similarly, the balance between spin relaxation and spin generation is expressed by

$$rsp + s/\tau_s = P_n(t = 0)G, \quad (10)$$

where $s = n_+ - n_-$ is the electron spin density and $P_n(t = 0)$ is the spin polarization at the moment of photoexcitation, given by Eq. (5). Holes are assumed to lose their spin orientation very fast, so they are treated as unpolarized. The first term in Eq. (10) describes the disappearance of the spin density due to carrier recombination, while the second term describes the intrinsic spin relaxation. From Eqs. (9) and (10) we obtain the steady-state electron polarization as (Žutić *et al.*, 2001b)

$$P_n = P_n(t = 0) \frac{1 - n_0 p_0 / np}{1 + 1/\tau_s rp}. \quad (11)$$

In a p-doped sample $p \approx p_0$, $n \gg n_0$, and Eq. (11) gives

$$P_n = P_n(t = 0)/(1 + \tau/\tau_s), \quad (12)$$

where $\tau = 1/rp_0$ is the electron lifetime.²⁹ The steady-state polarization is independent of the illumination intensity, being reduced from the initial spin polarization $P_n(t = 0)$.³⁰ The polarization of the photoluminescence is $P_{\text{circ}} = P_n(t = 0)P_n$ (Parsons, 1969). Early measurements of $P_n = 0.42 \pm 0.08$ in GaSb (Parsons, 1969) and $P_n = 0.46 \pm 0.06$ in Ga_{0.7}Al_{0.3}As (Ekimov and Safarov, 1970) showed an effective spin orientation close to the maximum value of $P_n(t = 0) = 1/2$ for a bulk unstrained zincblende structure, indicating that $\tau/\tau_s \ll 1$.

For spin pumping in an n-doped sample, where $n \approx n_0$ and $p \gg p_0$, Eqs. (9) and (11) give (D'yakonov and Perel', 1971b)

$$P_n = P_n(t = 0)/(1 + n_0/G\tau_s). \quad (13)$$

In contrast to the previous case, the carrier (now hole) lifetime $\tau = 1/rn_0$ has no effect on P_n . However, P_n depends on the photoexcitation intensity G , as expected for a pumping process. The effective carrier lifetime is $\tau_J = n_0/G$, where J represents the intensity of the illuminating light. If it is comparable to or shorter than τ_s , spin pumping is very effective. Spin pumping works because the photoexcited spin-polarized electrons do not

²⁹ After the illumination is switched off, the electron spin density, or equivalently the nonequilibrium magnetization, will decrease exponentially with the inverse time constant $1/T_s = 1/\tau + 1/\tau_s$ (Parsons, 1969).

³⁰ The effect of a finite length for the light absorption on P_n is discussed by Pierce and Celotta (1984). The absorption length α^{-1} is typically a micron for GaAs. It varies with frequency roughly as $\alpha(\hbar\omega) \propto (\hbar\omega - E_g)^{1/2}$ (Pankove, 1971).

need to recombine with holes. There are plenty of unpolarized electrons in the conduction band available for recombination. The spin is thus pumped in to the electron system.

When magnetic field \mathbf{B} is applied perpendicular to the axis of spin orientation (transverse magnetic field), it will induce spin precession with the Larmor frequency $\Omega_L = \mu_B g B / \hbar$, where μ_B is the Bohr magneton and g is the electron g factor.³¹ The spin precession, together with the random character of carrier generation or diffusion, leads to the spin dephasing (see Sec. III.A.1). Consider spins excited by circularly polarized light (or by any means of spin injection) at a steady rate. In a steady state a balance between nonequilibrium spin generated and spin relaxation is maintained, resulting in a net magnetization. If a transverse magnetic field is applied, the decrease of the steady-state magnetization can have two sources: (a) spins which were excited at random time and (b) random diffusion of spins towards a detection region. Consequently, spins precess along the applied field acquiring random phases relative to those which were excited or have arrived at different times. As a result, the projection of the electron spin along the exciting beam will decrease with the increase of transverse magnetic field, leading to depolarization of the luminescence. This is also known as the Hanle effect (Hanle, 1924), in analogy to the depolarization of the resonance fluorescence of gases. The Hanle effect was first measured in semiconductors by Parsons (1969). The steady-state spin polarization of the precessing electron spin can be calculated by solving the Bloch-Torrey equations (Bloch, 1946; Torrey, 1956), Eqs. (52)–(54) describing the spin dynamics of diffusing carriers.

In p-doped semiconductors the Hanle curve shows a Lorentzian decrease of the polarization (Parsons, 1969), $P_n(B) = P_n(B=0)/(1 + \Omega_L T_s)^2$, where $P_n(B=0)$ is the polarization at $B=0$ from Eq. (12) and T_s^{-1} is the effective spin lifetime given by $1/T_s = 1/\tau + 1/\tau_s$; see footnote 26. Measurements of the Hanle curve in GaAlAs were used by Garbuзов *et al.* (1971) to separately determine both τ and τ_s at various temperatures. The theory of the Hanle effect in n-doped semiconductors was developed by D'yakonov and Perel' (1976) who showed the non-Lorentzian decay of the luminescence for the regimes both of low ($\tau_J/\tau_s \gg 1$) and high ($\tau_J/\tau_s \ll 1$) intensity of the exciting light. At high fields $P_n(B) \propto 1/B^{1/2}$, consistent with the experiments of Vekua *et al.* (1976) in $\text{Ga}_{0.8}\text{Al}_{0.2}\text{As}$, showing a Hanle curve different from the usual $P_n(B) \propto 1/B^2$ Lorentzian behavior (D'yakonov and Perel', 1984a). Recent findings on the Hanle effect in nonuniformly doped GaAs and reanalysis of some earlier studies are given by Dzihoev *et al.* (2003).

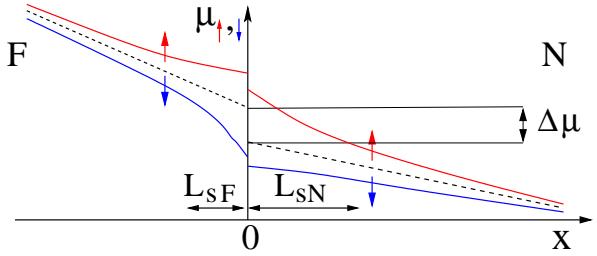


FIG. 7 Spatial variation of the electrochemical potentials near a spin-selective resistive interface at an F/N junction. At the interface $x = 0$ both the spin-resolved electrochemical potentials (μ_λ , $\lambda = \uparrow, \downarrow$, denoted with solid lines) and the average electrochemical potential (μ_F , μ_N , dashed lines) are discontinuous. The spin diffusion length L_{SF} and L_{SN} characterizes the decay of $\mu_s = \mu_\uparrow - \mu_\downarrow$ (or equivalently the decay of spin accumulation and the nonequilibrium magnetization) away from the interface and into the bulk F and N regions, respectively.

C. Theories of spin injection

Reviews on spin injection have covered materials ranging from semiconductors to high temperature superconductors and have addressed the implications for device operation as well as for fundamental studies in solid state systems.³² In addition to degenerate conductors, examined in these works, we also give results for nondegenerate semiconductors in which the violation of local charge neutrality, electric fields, and carrier band bending require solving the Poisson equation. The notation introduced here emphasizes the importance of different (and inequivalent) spin polarizations arising in spin injection.

1. F/N junction

A theory of spin injection across a ferromagnet/normal metal (F/N) interface was first offered by Aronov (1976b). Early work also included spin injection into a semiconductor (Sm) (Aronov and Pikus, 1976; Masterov and Makovskii, 1979) and a superconductor (S) (Aronov, 1976a). Spin injection in F/N junctions was subsequently studied in detail by Johnson and Silsbee (1987, 1988a),³³ van Son *et al.* (1987), Valet and Fert (1993), Hershfield and Zhao (1997), and others. Here we follow the approach of Rashba (2000, 2002c) and consider a steady-state³⁴ flow of electrons along the x direction in

³¹ In our convention the g factor of free electrons is positive, $g_0 = 2.0023$ (Kittel, 1996).

³² See, for example, (Goldman *et al.*, 2001, 1999; Jedema *et al.*, 2002d; Johnson, 2001, 2002a; Maekawa *et al.*, 2001; Osofsky, 2000; Schmidt and Molenkamp, 2002; Tang *et al.*, 2002; Wei, 2002)

³³ Johnson and Silsbee base their approach on irreversible thermodynamics and consider also the effects of a temperature gradient on spin-polarized transport, omitted in this section.

³⁴ Even some dc spin injection experiments are actually performed at low (audio-frequency) bias. Generalization to ac spin injection, with a harmonic time dependence, was studied by Rashba

a three-dimensional (3D) geometry consisting of a metallic ferromagnet (region $x < 0$) and a paramagnetic metal or a degenerate semiconductor (region $x > 0$).

The two regions, F and N, form a contact at $x = 0$, as depicted in Fig. 7. The relative magnitudes of three characteristic resistances per unit area³⁵ determine the degree of current polarization injected into a nonmagnetic material. These are the contact resistance r_c and the two characteristic resistances r_N, r_F , each given by the ratio of the spin diffusion length and the effective bulk conductivity in the corresponding region. Two limiting cases correspond to the transparent limit, where $r_c \rightarrow 0$, and the low-transmission limit, where $r_c \gg r_N, r_F$.

Spin-resolved quantities are labeled by $\lambda = 1$ or \uparrow for spin up, $\lambda = -1$ or \downarrow for spin down along the chosen quantization axis. For a free electron, spin angular momentum and magnetic moment are in opposite directions, and what precisely is denoted by “spin up” varies in the literature (Jonker *et al.*, 2003b). Conventionally, in metallic systems (Gijs and Bauer, 1997; Tedrow and Meservey, 1973), spin up refers to carriers with majority spin. This means that the spin (angular momentum) of such carriers is antiparallel to the magnetization. Spin-resolved charge current (density) in a diffusive regime can be expressed as

$$j_\lambda = \sigma_\lambda \nabla \mu_\lambda, \quad (14)$$

where σ_λ is conductivity and the electrochemical potential is

$$\mu_\lambda = (qD_\lambda/\sigma_\lambda)\delta n_\lambda - \phi, \quad (15)$$

with q proton charge, D_λ diffusion coefficient, $\delta n_\lambda = n_\lambda - n_{\lambda 0}$ the change of electron density from the equilibrium value for spin λ , and ϕ the electric potential.³⁶

In the steady state the continuity equation is

$$\nabla j_\lambda = \lambda q \left[\frac{\delta n_\lambda}{\tau_{\lambda-\lambda'}} - \frac{\delta n_{-\lambda}}{\tau_{-\lambda\lambda'}} \right], \quad (16)$$

and $\tau_{\lambda\lambda'}$ is the average time for flipping a λ -spin to λ' -spin. For a degenerate conductor³⁷ the Einstein relation is

$$\sigma_\lambda = q^2 \mathcal{N}_\lambda D_\lambda, \quad (17)$$

where $\sigma = \sigma_\uparrow + \sigma_\downarrow$ and $\mathcal{N} = \mathcal{N}_\uparrow + \mathcal{N}_\downarrow$ is the density of states. Using a detailed balance $\mathcal{N}_\uparrow/\tau_{\uparrow\uparrow} = \mathcal{N}_\downarrow/\tau_{\downarrow\downarrow}$

(2002b).

³⁵ For this simple geometry various resistances have a common factor of the cross-sectional area, which can be factored out. This is no longer possible for a more complicated geometry (Takahashi and Maekawa, 2003).

³⁶ More generally, for a noncollinear magnetization, j_λ becomes a second-rank tensor (Johnson and Silsbee, 1988a; Margulis and Margulis, 1994; Stiles and Zangwill, 2002).

³⁷ In the nondegenerate case of Boltzmann statistics, the Einstein relation implies that the ratio of the diffusion coefficient and the mobility is $k_B T/q$.

(Hershfield and Zhao, 1997; Kravchenko, 2002) together with Eqs. (15) and (17), the continuity equation can be expressed as

$$\nabla j_\lambda = \lambda q^2 \frac{\mathcal{N}_\uparrow \mathcal{N}_\downarrow}{\mathcal{N}_\uparrow + \mathcal{N}_\downarrow} \frac{\mu_\lambda - \mu_{-\lambda}}{\tau_s}, \quad (18)$$

where $\tau_s = \tau_{\uparrow\uparrow} \tau_{\downarrow\downarrow}/(\tau_{\uparrow\uparrow} + \tau_{\downarrow\downarrow})$ is the spin relaxation time. Equation (18) implies the conservation of charge current $j = j_\uparrow + j_\downarrow = \text{const.}$, while the spin counterpart, the difference of the spin-polarized currents $j_s = j_\uparrow - j_\downarrow$ is position dependent. Other “spin quantities,” X_s , unless explicitly defined, are analogously expressed with the corresponding (spin) polarization given by $P_X = X_s/X$. For example, the current polarization³⁸ $P_j = j_s/j$, generally different from the density polarization $P_n = (n_\uparrow - n_\downarrow)/n$, is related to the conductivity polarization P_σ as

$$P_j = 2(\sigma_\uparrow \sigma_\downarrow / \sigma) \nabla \mu_s / j + P_\sigma \quad (19)$$

where $\mu_s = \mu_\uparrow - \mu_\downarrow$. In terms of the average electrochemical potential $\mu = (\mu_\uparrow + \mu_\downarrow)/2$, P_σ further satisfies

$$\nabla \mu = -P_\sigma \nabla \mu_s / 2 + j / \sigma. \quad (20)$$

From Eqs. (15) and (18) it follows that μ_s satisfies the diffusion equation (Hershfield and Zhao, 1997; Schmidt *et al.*, 2000; Valet and Fert, 1993; van Son *et al.*, 1987)

$$\nabla^2 \mu_s = \mu_s / L_s^2, \quad (21)$$

where the spin diffusion length is $L_s = (\overline{D}\tau_s)^{1/2}$ with the spin averaged diffusion coefficient $\overline{D} = (\sigma_\downarrow D_\uparrow + \sigma_\uparrow D_\downarrow)/\sigma = \mathcal{N}(\mathcal{N}_\downarrow/D_\uparrow + \mathcal{N}_\uparrow/D_\downarrow)^{-1}$. Using Eq. (15) and the local charge quasineutrality $\delta n_\uparrow + \delta n_\downarrow = 0$ shows that μ_s is proportional to the nonequilibrium spin density $\delta s = \delta n_\uparrow - \delta n_\downarrow$ ($s = s_0 + \delta s = n_\uparrow - n_\downarrow$)

$$\mu_s = \frac{1}{2q} \frac{\mathcal{N}_\uparrow + \mathcal{N}_\downarrow}{\mathcal{N}_\uparrow \mathcal{N}_\downarrow} \delta s. \quad (22)$$

Correspondingly, μ_s is often referred to as the (nonequilibrium) *spin accumulation*³⁹ and is used to explain the GMR effect in CPP structures (Gijs and Bauer, 1997; Hartman (Ed.), 2000; Hirota *et al.*, 2002; Johnson, 1991; Valet and Fert, 1993).

The preceding equations are simplified for the N region by noting that $\sigma_\lambda = \sigma/2$, $\sigma_s = 0$, and $D_\lambda = \overline{D}$. Quantities pertaining to a particular region are denoted by the index F or N.

Equation (21) has also been used to study the diffusive spin-polarized transport and spin accumulation in

³⁸ This is also referred to as a spin injection coefficient (Rashba, 2000, 2002c).

³⁹ Spin accumulation is also relevant to a number of physical phenomena outside the scope of this article, for example, to the tunneling rates in the quantum Hall regime (Chan *et al.*, 1999; MacDonald, 1999).

ferromagnet/superconductor structures (Jedema *et al.*, 1999). Some care is needed to establish the appropriate boundary conditions at the F/N interface. In the absence of spin-flip scattering⁴⁰ at the F/N interface (which can arise, for example, due to spin-orbit coupling or magnetic impurities) the spin current is continuous and thus $P_{jF}(0^-) = P_{jN}(0^+) \equiv P_j$ (omitting $x = 0^\pm$ for brevity, and superscripts \pm in other quantities). These boundary conditions were used by Aronov (1976b); Aronov and Pikus (1976) without relating P_j to the effect of the F/N contact or material parameters in the F region.

Unless the F/N contact is highly transparent, μ_λ is discontinuous across the interface (Hershfield and Zhao, 1997; Johnson and Silsbee, 1988c; Rashba, 2000; Valet and Fert, 1993) and the boundary condition is

$$j_\lambda(0) = \Sigma_\lambda [\mu_{\lambda N}(0) - \mu_{\lambda F}(0)], \quad (23)$$

where

$$\Sigma = \Sigma_\uparrow + \Sigma_\downarrow \quad (24)$$

is the contact conductivity. For a free-electron model $\Sigma_\uparrow \neq \Sigma_\downarrow$ can be simply inferred from the effect of the exchange energy, which would yield spin-dependent Fermi wave vectors and transmission coefficients. A microscopic determination of the corresponding contact resistance [see Eq. (27)] is complicated by the influence of disorder, surface roughness, and different scattering mechanisms and is usually obtained from model calculations (Schep *et al.*, 1997; Stiles and Penn, 2000). Continued work on the first-principles calculation of F/N interfaces (Erwin *et al.*, 2002; Stiles, 1996) is needed for a more detailed understanding of spin injection. From Eqs. (23) and (24) it follows that

$$\mu_{sN}(0) - \mu_{sF}(0) = 2r_c(P_j - P_\Sigma)j, \quad (25)$$

$$\mu_N(0) - \mu_F(0) = r_c(1 - P_\Sigma P_j)j, \quad (26)$$

where the effective contact resistance is

$$r_c = \Sigma / 4\Sigma_\uparrow \Sigma_\downarrow. \quad (27)$$

The decay of μ_s , away from the interface, is characterized by the corresponding spin diffusion length

$$\mu_{sF} = \mu_{sF}(0)e^{x/L_{sF}}, \quad \mu_{sN} = \mu_{sN}(0)e^{-x/L_{sN}}. \quad (28)$$

A nonzero value for $\mu_{sN}(0)$ implies the existence of nonequilibrium magnetization δM in the N region (for noninteracting electrons $q\mu_s = \mu_B\delta M/\chi$, where χ is the magnetic susceptibility). Such a δM , as a result of electrical spin injection, was proposed by Aronov and Pikus (1976) and first measured in metals by Johnson and Silsbee (1985).

⁴⁰ The effects of non-conserving interfacial scattering on spin injection were considered in (Fert and Lee, 1996; Rashba, 2002c; Valet and Fert, 1993).

By applying Eq. (19), separately, to the F and N regions, one can obtain the amplitude of spin accumulation in terms of the current and density of states spin polarization and the effective resistances r_F and r_N ,

$$\mu_{sF}(0) = 2r_F [P_j - P_{\sigma F}] j, \quad \mu_{sN}(0) = -2r_N P_j j, \quad (29)$$

where

$$r_N = L_{sN}/\sigma_N, \quad r_F = L_{sF}\sigma_F/(4\sigma_{\uparrow F}\sigma_{\downarrow F}). \quad (30)$$

From Eqs. (29) and (25) the current polarization can be obtained as

$$P_j = [r_c P_\Sigma + r_F P_{\sigma F}] / r_{FN}, \quad (31)$$

where $r_{FN} = r_F + r_c + r_N$ is the effective equilibrium resistance of the F/N junction. It is important to emphasize that a measured highly polarized current, representing an efficient spin injection, does not itself imply a large spin accumulation or a large density polarization, typically measured by optical techniques. In contrast to the derivation of P_j from Eq. (31), determining P_n requires using Poisson's equation or a condition of the local charge quasineutrality.⁴¹

It is useful to note⁴² that Eq. (31), written as Eq. (18) in (Rashba, 2000) can be mapped to Eq. (A11) from (Johnson and Silsbee, 1987), where it was first derived.⁴³ An equivalent form for P_j in Eq. (31) was obtained by Hershfield and Zhao (1997) and for $r_c = 0$ results from van Son *et al.* (1987) are recovered.

In contrast to normal metals (Johnson and Silsbee, 1985, 1988d) and superconductors, for which injection has been reported in both conventional (Johnson, 1994), and high temperature superconductors (Dong *et al.*, 1997; Hass *et al.*, 1994; Vas'ko *et al.*, 1997; Yeh *et al.*, 1999), creating a substantial current polarization by direct electrical spin injection from a metallic ferromagnet into a semiconductor proved to be more difficult (Filip *et al.*, 2000; Hammar *et al.*, 1999; Monzon and Roukes, 1999; Zhu *et al.*, 2001).

By examining Eq. (31) we can both infer some possible limitations and deduce several experimental strategies for effective spin injection i.e. to increase P_j into

⁴¹ Carrier density will also be influenced by the effect of screening, which changes with the dimensionality of the spin injection geometry (Korenblum and Rashba, 2002).

⁴² Rashba (2002a).

⁴³ The substitutions are $P_j \rightarrow \eta^*$, $P_\sigma \rightarrow p$, $P_\Sigma \rightarrow \eta$, $r_c \rightarrow [G(\xi - \eta^2)]^{-1}$, $r_N \rightarrow \delta_n/\sigma_n \zeta_n$, $r_F \rightarrow \delta_f/\sigma_f (\zeta_f - p_f^2)$, $L_{sN,F} \rightarrow \delta_{n,f}$, and n, f label N and F region, respectively. η , ζ_n , and ζ_f are of the order of unity. To ensure that resistances and the spin diffusion lengths in Johnson and Silsbee (1987) are positive, one must additionally have $(\xi - \eta^2) > 0$ and $(\zeta_i - p_i^2) > 0$, $i = n, f$ (for normal and ferromagnetic regions, respectively). In particular, assuming $\xi = \zeta_n = \zeta_f = 1$ a detailed correspondence between Eq. (31) and Eq. (A11) in (Johnson and Silsbee, 1987) is recovered. For example, $r_c \rightarrow [G(\xi - \eta^2)]^{-1}$ yields Eq. (27), where $\Sigma \rightarrow G$.

semiconductors. For a perfect Ohmic contact $r_c = 0$, the typical resistance mismatch $r_F \ll r_N$ (where F is a metallic ferromagnet) implies inefficient spin injection with $P_j \approx r_F/r_N \ll 1$, referred to as the *conductivity mismatch* problem by Schmidt *et al.* (2000). Even in the absence of the resistive contacts, effective spin injection into a semiconductor can be achieved if the resistance mismatch is reduced by using for spin injectors either a magnetic semiconductor or a highly spin-polarized ferromagnet.⁴⁴

While there was early experimental evidence (Alvarado and Renaud, 1992) that employing resistive (tunneling) contacts could lead to an efficient spin injection⁴⁵ a systematic understanding was provided by Rashba (2000) and supported with the subsequent experimental and theoretical studies (Fert and Jaffres, 2001; Johnson, 2003; Johnson and Byers, 2003; Rashba, 2002c; Smith and Silver, 2001; Takahashi and Maekawa, 2003). As can be seen from Eq. (31) the spin-selective resistive contact $r_c \gg r_F, r_N$ (such as a tunnel or Schottky contact) would contribute to effective spin injection with $P_j \approx P_\Sigma$ being dominated by the effect r_c and not the ratio r_F/r_N .⁴⁶ This limit is also instructive to illustrate the principle of spin filtering (Esaki *et al.*, 1967; Filip *et al.*, 2002; Hao *et al.*, 1990; Moodera *et al.*, 1988). In a spin-discriminating transport process the resulting degree of spin polarization is changed. Consequently the effect of spin filtering, similar to spin injection, leads to the generation of (nonequilibrium) spin polarization.⁴⁷ For example, at low temperature EuS and EuSe, discussed in Sec. IV.C, can act as spin-selective barriers. In the extreme case, initially spin-unpolarized carriers (say, injected from a nonmagnetic material) via spin-filtering could attain a complete polarization. For a strong spin-filtering contact $P_\Sigma > P_{\sigma F}$, the sign of the spin accumulation (nonequilibrium magnetization) is reversed in the F and N regions, near the interface [recall Eq. (25)], in contrast to the behavior sketched in Fig. 7, where $\mu_{sF,N} > 0$.

The spin injection process alters the potential drop across the F/N interface because differences of spin-dependent electrochemical potentials on either side of the interface generate an effective resistance δR . By integrating Eq. (20) for N and F regions, separately, it follows that $R_j = \mu_N(0) - \mu_F(0) + P_{\sigma F}\mu_{sF}(0)/2$, where R is the junction resistance. Using Eqs. (26), (30), and (31) allows us to express $R = R_0 + \delta R$, where $R_0 = 1/\Sigma$ ($R_0 = r_c$ if $\Sigma_\uparrow = \Sigma_\downarrow$) is the equilibrium resistance, in the

absence of spin injection, and

$$\delta R = [r_N(r_F P_{\sigma F}^2 + r_c P_\Sigma^2) + r_F r_c (P_{\sigma F} - P_\Sigma)^2]/r_{FN}, \quad (32)$$

where $\delta R > 0$ is the nonequilibrium resistance. Petukhov has shown (Jonker *et al.*, 2003a) that Eqs. (31) and (32) could be obtained by considering an equivalent circuit scheme with two resistors $\tilde{R}_\uparrow, \tilde{R}_\downarrow$ connected in parallel, where $\tilde{R}_\lambda = L_{sF}/\sigma_{\lambda F} + 1/\Sigma_\lambda + 2L_{sN}/\sigma_N$ and $\tilde{R}_\uparrow + \tilde{R}_\downarrow = 4r_{FN}$. For such a resistor scheme, by noting that $j_\uparrow \tilde{R}_\uparrow = j_\downarrow \tilde{R}_\downarrow$, Eq. (31) is obtained as $P_j = -P_{\tilde{R}} \equiv -(\tilde{R}_\uparrow - \tilde{R}_\downarrow)/(\tilde{R}_\uparrow + \tilde{R}_\downarrow)$. δR in Eq. (32) is then obtained as the difference between the total resistance of the nonequilibrium spin-accumulation region of the length $L_{sF} + L_{sN}$ [given by the equivalent resistance $\tilde{R}_\uparrow \tilde{R}_\downarrow / (\tilde{R}_\uparrow + \tilde{R}_\downarrow)$] and the equilibrium resistance for the same region, $L_{sF}/\sigma_F + L_{sN}/\sigma_N$.

The concept of the excess resistance δR can also be explained as a consequence of the Silsbee-Johnson spin-charge coupling Johnson and Silsbee (1985, 1987); Silsbee (1980) and illustrated by considering the simplified schemes in Figs. 5 and 7. Accumulated spin near the F/N interface, together with a finite spin relaxation and a finite spin diffusion, impedes the flow of spins and acts as a “spin bottleneck” (Johnson, 1991). A rise of μ_{sN} must be accompanied by the rise of μ_{sF} [their precise alignment at the interface is given in Eq. (25)] or there will be a backflow of the nonequilibrium spin back into the F region. Because both spin and charge are carried by electrons in spin-charge coupling the backflow of spin driven by diffusion creates an additional resistance for the charge flow across the F/N interface. Based on an analogy with the charge transport across a clean N/superconductor (S) interface (see Sec. IV.A.3) van Son *et al.* (1987) explained δR by invoking the consequences of current conversion from spin-polarized, at far to the left of the F/N interface, to completely unpolarized, at far right in the N region.

The increase in the total resistance with spin injection can be most dramatic if the N region is taken to be a superconductor (S); see Fig. 5(c). Spin injection depletes the superconducting condensate and can result in the switching to a normal state of much higher resistance (Dong *et al.*, 1997; Takahashi *et al.*, 1999; Vas'ko *et al.*, 1997; Wei *et al.*, 1999; Yeh *et al.*, 1999). A critical review of possible spurious effects in reported experiments Gim *et al.* (2001) has also stimulated the development of a novel detection technique which uses scanning tunneling spectroscopy combined with pulsed quasiparticle spin injection to minimize Joule heating (Ngai *et al.*, 2004) (see Sec. IV.A.1). In the S region the quasiparticle energy is $E_k = (\xi_k^2 + \Delta^2)^{1/2}$, where ξ_k is the single particle excitation energy corresponding to the wave vector \mathbf{k} and Δ is the superconducting gap [see Fig. 5(c)]. Such a dispersion relation results in a smaller diffusion coefficient and a longer spin flip time than in the N region, while their product, the spin diffusion length, remains the same (Yamashita *et al.*, 2002). Consequently, Eq. (21) also

⁴⁴ From Eq. (30) a half-metallic ferromagnet implies a large r_F .

⁴⁵ The influence of the resistive contacts on spin injection can also be inferred by explicitly considering resistive contacts (Hershfield and Zhao, 1997; Johnson and Silsbee, 1987).

⁴⁶ A similar result was stated previously by Johnson and Silsbee (1988a).

⁴⁷ While most of the schemes resemble a CPP geometry [Fig. 3(b)], there are also proposals for generating highly polarized currents in a CIP-like geometry [Fig. 3(a)] (Gurzhi *et al.*, 2001, 2003).

applies to the diffusive spin-polarized transport and spin accumulation in ferromagnet/superconductor structures (Jedema *et al.*, 1999; Yamashita *et al.*, 2002). Opening of the superconducting gap implies that a superconductor is a low carrier system for spin, which is carried by quasiparticles (Takahashi and Maekawa, 2003).

In the preceding analysis, appropriate for bulk, homogeneous, three-dimensional N and F regions and degenerate (semi)conductors, Poisson's equation was not invoked and the local charge neutrality $\delta n_\uparrow + \delta n_\downarrow$ was used only to derive Eq. (22).⁴⁸ Focusing on bulk samples in which both the size of the F and N regions and the corresponding spin diffusion lengths are much larger than the Debye screening length, one can find that the quasineutrality condition, combined with the Eqs. (15) and (17), yields

$$\phi = -\mu - P_N \mu_s / 2, \quad (33)$$

where the density of states spin polarization of P_N vanishes in the N region. At the contact $x = 0$ there is a potential drop, even when $r_c = 0$, which can be evaluated from Eqs. (26) and (33) as

$$\phi_N(0) - \phi_F(0) = -r_c [1 - P_\Sigma P_j] j + P_{NF}(0) \mu_{sF}(0) / 2. \quad (34)$$

The creation of nonequilibrium spin in the N region results in the spin EMF in the F/N structure which can be used to detect electrical spin injection, as depicted in Fig. 5. Within a simplified semi-infinite geometry for the F and N regions, we consider an effect of spin pumping in the N region, realized either by electrical spin injection from another F region [as shown in Fig. 5(b)] or by optical pumping (see Sec. II.B). The resulting potential drop can be calculated by modifying μ_{sN} in Eq. (28),

$$\mu_{sN} = \mu_{sN}(\infty) + [\mu_{sN}(0) - \mu_{sN}(\infty)] e^{-x/L_{sN}}, \quad (35)$$

where $\mu_{sN}(\infty)$ represents the effect of homogeneous spin pumping in the N region. To calculate the open circuit voltage ($j = 0$) the continuity of spin current at $x = 0$ should be combined with the fact that $P_j j = j_s$. From Eq. (19) it follows that

$$j_s(0) = 2 \frac{\sigma_\uparrow \sigma_\downarrow}{\sigma_F} \frac{\mu_{sF}(0)}{L_{sF}} = -\frac{1}{2} \sigma_N \frac{\mu_{sN}(0) - \mu_{sN}(\infty)}{L_{sN}}, \quad (36)$$

while the discontinuity of μ_s in Eq. (25) yields⁴⁹

$$\begin{aligned} \mu_{sF}(0) &= (r_F/r_{FN}) \mu_{sN}(\infty), & j_s(0) &= \mu_{sN}(\infty) / 2r_{FN}, \\ \mu_{sN}(0) &= [(r_c + r_F)/r_{FN}] \mu_{sN}(\infty). \end{aligned} \quad (37)$$

By substituting this solution into Eq. (34), we can evaluate the contact potential drop can be evaluated as

$$\phi_N(0) - \phi_F(0) = [r_F P_{NF} + r_c P_\Sigma] \mu_{sN}(\infty) / 2r_{FN}. \quad (38)$$

The total potential drop (recall $j = 0$) at the F/N junction⁵⁰ is (Rashba, 2002c)

$$\Delta\phi_{FN} = \phi_N(\infty) - \phi_F(-\infty) = P_j \mu_{sN}(\infty) / 2. \quad (39)$$

where P_j is given in Eq. (31). In the context of the spin-detection scheme from Fig. 5 and high impedance measurements at the N/F2 junction, the spin-coupled voltage V_s (Johnson and Silsbee, 1985; Silsbee, 1980) was also found to be proportional to current polarization and the spin accumulation ($\mu_s \propto \delta s \propto \delta M$) (Johnson and Silsbee, 1988b).

2. F/N/F junction

The above analysis of the F/N bilayer can be readily extended to the geometry in which the infinite F regions are separated by an N region of thickness d . The quantities pertaining to the two ferromagnets are defined as in the case of an F/N junction and labeled by the superscripts L and R (left and right regions, respectively). It follows from Eq. (19), by assuming the continuity of the spin current at L,R, that the difference of the spin-resolved electrochemical potential, responsible for the spin accumulation, is

$$\mu_{sF}^L = 2r_F^L (P_j^L - P_{\sigma F}^L) j e^{x/L_{sF}^L}, \quad x < 0, \quad (40)$$

$$\begin{aligned} \mu_{sN} &= 2r_N [P_j^R \cosh(x/L_{sN}) - P_j^L \cosh[(d-x)/L_{sN}]] \\ &\times j / \sinh(d/L_{sN}), \quad 0 < x < d, \end{aligned} \quad (41)$$

$$\mu_{sF}^R = -2r_F^R (P_j^R - P_{\sigma F}^R) j e^{(d-x)/L_{sF}^R}, \quad x > d, \quad (42)$$

where the current spin polarization $P_j^{L,R}$ at the two contacts in the F/N/F geometry can be expressed (Rashba, 2002c) in terms of the P_j calculated for F/N junction with the infinite F and N regions in Eq. (19) and the appropriate effective resistances. By $P_{j\infty}^{L,R}$ we denote the P_j calculated in Eq. (31) for at left and right contact (with the appropriate parameters for the F/N/F junction) as if it were surrounded by the infinite F and N regions. Analogously to the F/N junction, the consequence of the spin injection is the increase of the resistance $R = R_0 + \delta R$, as compared to the equilibrium value $R_0 = (\Sigma^L)^{-1} + (\Sigma^R)^{-1}$. The nonequilibrium resistance δR is also always positive for spin-conserving contacts (Rashba, 2000, 2002c), in agreement with experiments on all-semiconductor trilayer structures (Schmidt *et al.*, 2001); see Sec. II.D.3.

Many applications based on magnetic multilayers rely on the spin-valve effect in which the resistance changes due to the relative orientations of the magnetization in the two F regions. The geometry considered here is relevant for CPP GMR (Bass and Pratt, Jr., 1999; Gijs and

⁴⁸ For spin injection in nondegenerate semiconductors (with the carriers obeying the Boltzmann statistics) there can be large effects due to built-in fields and deviation from local charge neutrality, as discussed in Sec. II.C.3.

⁴⁹ A missprint in $\mu_{sF}(0)$ from Rashba (2002c) has been corrected.

⁵⁰ A similar potential drop was also calculated across a ferromagnetic domain wall (Dzero *et al.*, 2003).

Bauer, 1997; Parkin, 2002) and all-metallic spin injection of Johnson and Silsbee (1985). In particular, the resistance change between antiparallel and parallel magnetization orientations in the two ferromagnets can be expressed using current polarization of an infinite F/N junction $P_{j\infty}^{L,R}$ (Rashba, 2002c):

$$\Delta R = R_{\uparrow\downarrow} - R_{\uparrow\uparrow} = 4P_{j\infty}^L P_{j\infty}^R \frac{r_F^L r_{FN}^R r_N}{\mathcal{D} \sinh(d/L_{sN})}, \quad (43)$$

where $r_F^{L,R}$, $r_c^{L,R}$, and r_N are defined as in the case of an F/N junction and

$$\begin{aligned} \mathcal{D} &= (r_F^L + r_c^L)(r_c^R + r_F^R) \\ &+ r_N^2 + r_N(r_F^L + r_c^L + r_c^R + r_F^R) \coth(d/L_{sN}). \end{aligned} \quad (44)$$

Up to a factor of 2, Eq. (43) has also been obtained by Hershfield and Zhao (1997) using Onsager relations. In the limit of a thin N region, $d/L_{sN} \rightarrow 0$, ΔR remains finite. In the opposite limit, for $d \gg L_{sN}$,

$$\Delta R \sim P_{j\infty}^L P_{j\infty}^R \exp(-d/L_{sN}). \quad (45)$$

For a symmetric F/N/F junction, where $r_{c,F}^L = r_{c,F}^R$, it follows that

$$\Delta R = \frac{4r_N(r_c P_\Sigma + r_F P_\sigma F)}{\mathcal{D} \sinh(d/L_{sN})}. \quad (46)$$

Considering the spin injection from F into a ballistic N region in the presence of diffusive interfacial scattering, where the phase coherence is lost and the Boltzmann equation can be applied, it is instructive to reconsider the effect of contact resistance (Kravchenko and Rashba, 2003). We introduce the Sharvin resistance R_{Sharvin} (Sharvin, 1965), arising in ballistic transport between the two infinite regions connected by a contact (an orifice or a narrow and short constriction) of radius much smaller than the mean free path, $a \ll l$. In a 3D geometry the resistance is

$$R_{\text{Sharvin}} = \frac{4\rho l}{3\pi a^2} = \left[\frac{e^2 k^2 A}{h 2\pi} \right]^{-1}, \quad (47)$$

where $h/e^2 \approx 25.81 \text{ k}\Omega$ is the quantum of resistance per spin, A is the contact area, and k is the Fermi wave vector. The opposite limit, of diffusive transport through the contact with $a \gg l$, corresponds to the the Maxwell or Drude resistance $R_{\text{Maxwel}} = \rho/2a$. The studies of intermediate cases provide an interpolation scheme between the R_{Maxwell} and R_{Sharvin} for various ratios of a/l (de Jong, 1994; Jansen *et al.*, 1980; Nikolić and Allen, 1999; Wexler, 1966). Following Kravchenko and Rashba (2003) the effective contact resistance $r_c = r_{c\uparrow} + r_{c\downarrow}$ (recall that it is defined per unit area) is obtained as

$$r_{c\lambda} = (4R_{\text{Sharvin}}/A)(1 - t_\lambda^L - t_\lambda^R)/t_\lambda^L, \quad (48)$$

where $t_\alpha^{L,R}$ represent the transmission coefficients for the electrons reaching the contact from the left and from

the right and satisfy $t^L + t^R \leq 1$. For r_c which would exceed the resistance of the N and F bulk regions the spin injection efficiency can attain $P_j \sim (r_{c\uparrow} - r_{c\downarrow})/r_c$ (Kravchenko and Rashba, 2003), showing, similarly to the diffusive regime, the importance of the resistive contacts to efficient spin injection. Connection with the results in the diffusive regime can be obtained (Kravchenko and Rashba, 2003) by identifying $r_{c\lambda} = 1/4\Sigma_\lambda$, where the contact conductivity Σ_λ was introduced in Eq. 24.

While most of the experimental results on spin injection are feasible in the diffusive regime, there are many theoretical studies treating the ballistic case and phase-coherent transport both in F/N and F/N/F junctions (Hu and Matsuyama, 2001; Hu *et al.*, 2001a; Matsuyama *et al.*, 2002; Mireles and Kirczenow, 2001). Simple models in which the N region is a degenerate semiconductor often adopt an approach developed first for charge transport in junctions involving superconductors, discussed in Sec. IV.A.3. Considering spin-orbit coupling and the potential scattering at the F/N interface modeled by the δ -function, Hu and Matsuyama (2001) have examined ballistic spin injection in the F/N junction. They show that even a spin-independent barrier can be used to enhance the spin injection and lead to an increase in the conductance polarization. First-principles calculations were also used for ballistic spin injection from a ferromagnetic metal into a semiconductor (Mavropoulos *et al.*, 2002; Wunnicke *et al.*, 2002; Zwierzycki *et al.*, 2003). In the limit of coherent (specular) scattering⁵¹ and high interfacial quality it was shown that different band structure in the F and the N regions would contribute to a significant contact resistance and an efficient spin injection (Zwierzycki *et al.*, 2003).

3. Spin injection through the space-charge region

Interfaces making up a semiconductor often develop a space-charge region—a region of local macroscopic charges. Typical examples are the Schottky contact and the depletion layer in *p-n* junctions. While phenomenological models, such as the one introduced in Sec. II.C.1, capture a remarkable wealth of spin injection physics, they carry little information about spin-dependent processes right at the interfaces. Microscopic studies of spin-polarized transport and spin-resolved tunneling through space-charge regions are still limited in scope. The difficulty lies in the need to consider self-consistently simultaneous charge accumulation and electric-field generation (through Poisson's equation), both affecting transport. Non-self-consistent analyses of a Schottky barrier spin injection were performed in (Albrecht and Smith, 2002, 2003; Prins *et al.*, 1995), while Osipov and Bratkovsky

⁵¹ The wave-vector component along the interface is conserved during scattering.

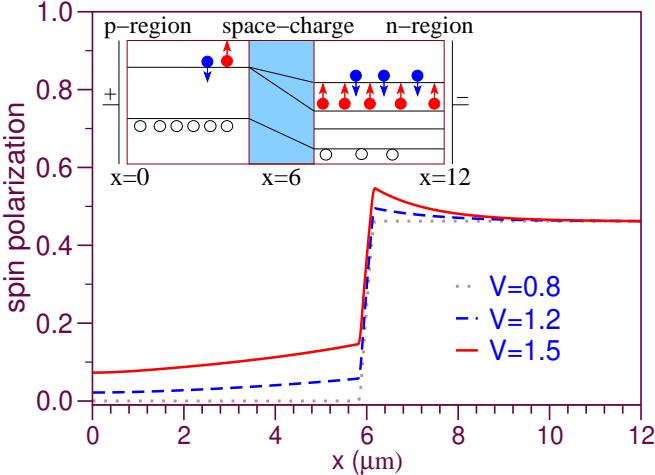


FIG. 8 Spin injection through the space-charge region of a magnetic p - n junction. The geometry is depicted in the inset, which shows a junction with a spin-split conduction band in the n -region with spin-polarized electrons (solid circles) and unpolarized holes (empty circles). Under applied forward bias V the charge current flows to the right. The curves, labeled by V , show the electron density polarization profiles $P_n(x)$ for the depicted geometry and GaAs materials parameters. The equilibrium density polarization in the n -region is about 0.5. At low bias (0.8 V) there is no spin injection. Spin injection, manifested by the increase of P_n in the p -region, appears only at large biases (1.2 and 1.5 V), where it is driven by electric drift (Žutić *et al.*, 2002). Spin polarization of the current is discussed by Fabian *et al.* (2002b); Žutić *et al.* (2001a). Adapted from Žutić *et al.*, 2002.

(2003) proposed an efficient spin injection method using a δ -doped Schottky contact.

Let us now consider spin injection through the depletion layer in magnetic p - n junctions (Fabian *et al.*, 2002b; Žutić *et al.*, 2002, 2003). The physics is based on drift and diffusion⁵² limited by carrier recombination and spin relaxation, as described in more detail in Sec. IV.A.4. The transport equations are solved self-consistently with Poisson's equation, taking full account of electric field due to accumulated charges. Additional examples of magnetic p - n junctions are discussed in Sec. IV.D.

The system is depicted in Fig. 8. The p - n junction has a magnetic n -region⁵³ with a net equilibrium electron spin P_{n0}^R , where R stands for the right (here n) region. Holes are assumed to be unpolarized. An important issue to be resolved is whether there will be spin accumulation

in the p -region if a forward bias is applied to the junction. In other words, will spin be injected across the depletion layer? Naively the answer is yes, since spin is carried by electrons, but the result shown in Fig. 8 suggests a more complicated answer. At small biases there is no spin injection. This is the normal limit of diode operation, in which where the injected carrier density through the depletion region is still smaller than the equilibrium carrier density. Only with bias increasing to the high-injection limit (typically above 1 V) is spin injected.

The explanation for the absence of spin injection at small biases and for nondegenerate doping levels (Boltzmann statistics are applicable) is as follows. On the n side, there are more spin up than spin down electrons, $n_\uparrow > n_\downarrow$. If $2q\zeta$ is the spin splitting of the conduction band, $n_\uparrow(\zeta)/n_\downarrow(\zeta = 0) = \exp(q\zeta/k_B T)$. Under a forward bias, electrons flow to the p -region. The flow is limited by thermal activation over the barrier (given by the built-in electrostatic potential minus bias), which is, for the spin up electrons, greater by $q\zeta$. For Boltzmann statistics, the rate of transmission of spin up electrons over the barrier is $\sim \exp(-q\zeta/k_B T)$. Since current is proportional to both the carrier density and the transmission rate, the two exponential factors cancel out. Similarly for spin down. As a result, the spin-resolved current is unaffected by $2q\zeta$ and there is no spin current flowing through the depletion layer. There is no spin accumulation. Spin injection appears only at large biases, where it is driven by electric drift leading to nonequilibrium spin population already in the n -region (Fabian *et al.*, 2002b; Žutić *et al.*, 2002). In addition to spin injection, spin extraction has also been predicted in magnetic p - n junctions with a magnetic p -region (Žutić *et al.*, 2002). Under a large bias, spin is extracted (depleted) from the nonmagnetic n -region.

Electric field in the bulk regions next to the space charge is important only at large biases. It affects not only spin density, but spin diffusion as well. That spin injection efficiency can increase in the presence of large electric fields due to an increase in the spin diffusion length (spin drag) was first shown by Aronov and Pikus (1976), and was later revisited by other authors.⁵⁴ To be important, the electric field needs to be very large,⁵⁵ more than 100 V/cm at room temperature. While such large fields are usually present inside the space-charge regions, they exist in the adjacent bulk regions only at the high injection limit and affect transport and spin injection. In addition to electric drift, magnetic drift,

⁵² Tunneling or field emission becomes important, for example, in thin Schottky barriers or in p - n junctions and heterostructures at large reverse biases (Johnston-Halperin *et al.*, 2002; Kohda *et al.*, 2001; Van Dorpe *et al.*, 2003a).

⁵³ Equilibrium magnetization can be a consequence of doping with magnetic impurities, yielding large carrier g factors, and applying magnetic field, or of using a ferromagnetic semiconductor (Ohno, 1998; Pearton *et al.*, 2003).

⁵⁴ See, for example, (Bratkovsky and Osipov, 2003; Fabian *et al.*, 2002b; Flensberg *et al.*, 2001; Margulis and Margulis, 1994; Martin, 2003; Vignale and D'Amico, 2003; Yu and Flatté, 2002a; Žutić *et al.*, 2001b).

⁵⁵ The critical magnitude is obtained by dividing a typical energy, such as the thermal or Fermi energy, by q and by the spin diffusion length. At room temperature the thermal energy is 25 meV, while the spin diffusion length can be several microns.

in magnetically inhomogeneous semiconductors, can also enhance spin injection (Fabian *et al.*, 2002b).

The following formula was obtained for spin injection at small biases (Fabian *et al.*, 2002b):

$$P_n^L = \frac{P_{n0}^L [1 - (P_{n0}^R)^2] + \delta P_n^R (1 - P_{n0}^L P_{n0}^R)}{1 - (P_{n0}^R)^2 + \delta P_n^R (P_{n0}^L - P_{n0}^R)}, \quad (49)$$

where L (left) and R (right) label the edges of the space-charge (depletion) region of a p - n junction. Correspondingly, δP_n^R represents the nonequilibrium electron polarization, evaluated at R , arising from a spin source. The case discussed in Fig. 8 is for $P_{n0}^L = \delta P_n^R = 0$. Then $P_n^L = 0$, in accord with the result of no spin injection. For a homogeneous equilibrium magnetization ($P_{n0}^L = P_{n0}^R$), $\delta P_n^L = \delta P_n^R$; the nonequilibrium spin polarization is the same across the depletion layer. Equation (49) demonstrates that only *nonequilibrium* spin, already present in the bulk region, can be transferred through the depletion layer at small biases (Fabian *et al.*, 2002b; Žutić *et al.*, 2001b). Spin injection of nonequilibrium spin is also very effective if it proceeds from the p -region (Žutić *et al.*, 2001b), which is the case for a spin-polarized solar cell (Žutić *et al.*, 2001a). The resulting spin accumulation in the n -region extends the spin diffusion range, leading to spin amplification—increase of the spin population away from the spin source. These results were also confirmed in the junctions with two differently doped n -regions (Pershin and Privman, 2003a,b). Note, however, that the term “spin polarization density” used in Pershin and Privman (2003a,b) is actually the spin density $s = n_\uparrow - n_\downarrow$, not the spin polarization P_n .

Theoretical understanding of spin injection has focused largely on spin density while neglecting spin phase, which is important for some proposed spintronic applications. The problem of spin evolution in various transport modes (diffusion, tunneling, thermionic emission) remains to be investigated. Particularly relevant is the question of whether spin phase is conserved during spin injection. Malajovich *et al.* (2001) showed, by studying spin evolution in transport through a n-GaAs/n-ZnSe heterostructure, that the phase can indeed be preserved.

D. Experiments on spin injection

1. Johnson-Silsbee spin injection

The first spin polarization of electrons by electrical spin injection (Johnson and Silsbee, 1985) was demonstrated in a “bulk wire” of aluminum on which an array of thin film permalloy (Py) pads (with 70 % nickel and 30 % iron) was deposited spaced in multiples of 50 μm , center to center (Johnson and Silsbee, 1988d) to serve as spin injectors and detectors. In one detection scheme a single ferromagnetic pad was used as a spin injector while the distance to the spin detector was altered by selecting different Py pads to detect V_s and through the spatial decay

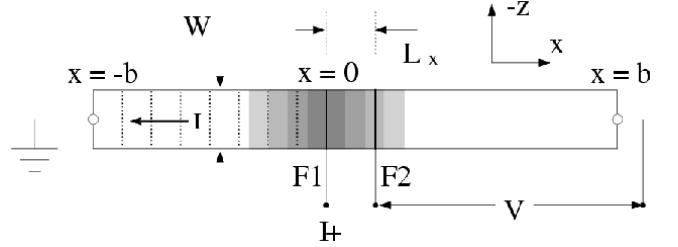


FIG. 9 Schematic top view of nonlocal, quasi-one-dimensional geometry used by Johnson and Silsbee (1985): F1 and F2, the two metallic ferromagnets having magnetizations in the x - z plane; dotted lines, equipotentials characterizing electrical current flow; grey shading, diffusing population of nonequilibrium spin-polarized electrons injected at $x = 0$, with darker shades corresponding to higher density of polarized electrons. From Johnson, 2002a.

of this spin-coupled voltage infer L_{sN} .⁵⁶ This procedure is illustrated in Fig. 9, where the separation between the spin injector and detector L_x is variable.

Johnson and Silsbee (1985) point out that in the depicted geometry there is no flow of the charge current for $x > 0$ and that in the absence of nonequilibrium spins a voltage measurement between $x = L_x$ and $x = b$ gives zero. Injected spin-polarized electrons will diffuse symmetrically (at low current density the effect of electric fields can be neglected), and the measurement of voltage will give a spin-coupled signal V_s related to the relative orientation of magnetizations in F1 and F2.⁵⁷ The results, corresponding to the polarizer-analyzer detection and the geometry of Fig. 9, are given in Fig. 10. An in-plane field ($\mathbf{B} \parallel \hat{\mathbf{z}}$), of a magnitude several times larger than a typical field for magnetization reversal, $B_0 \approx 100$ G, is applied to define the direction of magnetization in the injector and detector. As the field sweep is performed, from negative to positive values, at B_{01} there is a reversal of magnetization in one of the ferromagnetic films accompanied by a sign change in the spin-coupled signal. As B_z is further increased, at approximately B_{02} , there is another reversal of magnetization, resulting in parallel orientation of F1 and F2 and a V_s of magnitude similar to that for the previous parallel orientation when $B_z < B_{01}$.

A more effective detection of the spin injection is realized through measurements of the Hanle effect, also discussed Secs. II.B and III.A.2, and described by the Bloch-Torrey equations (Bloch, 1946; Torrey, 1956) [see Eqs. (52)–(54)]. The inset of Fig. 10 summarizes results

⁵⁶ The spin relaxation time in a ferromagnet is often assumed to be very short. Correspondingly, in the analysis of the experimental data, both the spin diffusion length and δM are taken to vanish in the F region (Johnson and Silsbee, 1985, 1988a,d; Silsbee, 1980).

⁵⁷ This method for detecting the effects of spin injection is also referred to as a potentiometric method.

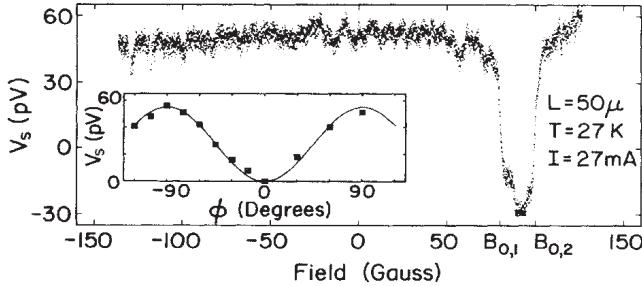


FIG. 10 Spin injection data from bulk Al wire sample. Negative magnetic field is applied parallel to the magnetization ($-z$ axis) in the two ferromagnetic regions. As the field is increased, at $B_{0,1}$ magnetization in one of the ferromagnetic regions is reversed, and at $B_{0,2}$ the magnetization in the other region is also reversed (both are along $+z$ axis). Inset: amplitude of the observed Hanle signal as a function of orientation angle ϕ of magnetic field. From Johnson and Silsbee, 1985.

from a series of Hanle experiments on a single sample. For the Hanle effect \mathbf{B} must have a component perpendicular to the orientation axes of the injected spins. Only projection of \mathbf{B} perpendicular to the spin axis applies a torque and dephases spins. The magnitude of \mathbf{B} , applied at an angle ϕ to the z -axis in the $y - z$ plane, is small enough that the magnetizations in ferromagnetic thin films remain in the $x - z$ plane (see Fig. 9). If, at $\mathbf{B} = 0$, injected nonequilibrium magnetization is $\delta M(0)\hat{z}$ then at finite field $\delta\mathbf{M}$ precesses about \mathbf{B} with the cone of angle 2ϕ . After averaging over several cycles, only $\delta M(0) \cos \phi$, the component $\parallel \mathbf{B}$, will survive. The voltage detector⁵⁸ senses the remaining part of the magnetization projected on the axis of the detector $\delta M(0) \cos \phi \times \cos \phi$ (Johnson and Silsbee, 1988a). The predicted angular dependence for the amplitude of the Hanle signal (proportional to the depolarization of δM in a finite field) $[\delta M(0) - \delta M(0) \cos^2 \phi]$ is plotted in the inset together with the measured data.⁵⁹ Results confirm the first application of the Hanle effect to dc spin injection.

The Hanle effect was also studied theoretically by solving the Bloch-Torrey equations for an arbitrary orientation, characterized by the angle α , between the magnetization in F1 and F2 (Johnson and Silsbee, 1988a). From the Hanle curve [$V_s(B_\perp)$] measured at $T = 4.3$ (36.6) K, the parameters $L_s = 450$ (180) μm and $P_\Sigma = 0.06$ (0.08) were extracted.⁶⁰ This spin injection technique using a few pV resolution of a superconducting quantum interfer-

ence device (SQUID) and with an estimated $P_\Sigma \approx 0.07$ provided an accuracy able to detect $P_n \approx 5 \times 10^{-12}$, causing speculating on that a single-spin sensitivity might be possible in smaller samples (Johnson and Silsbee, 1985, 1988d). While in a good conductor, such as Al, the observed resistance change ΔR was small ($\sim n\Omega$), the relative change at low temperatures and for $L_x \ll L_s$ was $\Delta R/R \approx 5\%$, where ΔR is defined as in Eq. (1), determined by the relative orientation of the magnetization in F1 and F2, and R is the Ohmic resistance (Johnson, 2002a). Analysis from Sec. II.C.2 shows that the measurement of ΔR could be used to determine the product of injected current polarizations in the two F/N junctions.

The studies of spin injection were extended to the thin-film geometry, also known as the “bipolar spin switch” or “Johnson spin transistor” (Johnson, 1993a,b) similar to the one depicted in Fig. 5(a). The measured spin-coupled signals⁶¹ in Au films were larger than the values obtained in bulk Al wires (Johnson and Silsbee, 1985, 1988d). A similar trend, $V_s \sim 1/d$, potentially important for applications, was already anticipated by Silsbee (1980). The saturation of this increase can be inferred from Eqs. (43) and (44) for $d \ll L_{sN}$ and has been discussed by Hershfield and Zhao (1997) and Fert and Lee (1996).

When polarizer-analyzer detection was used, one of the fitting parameters from the measured data P_Σ sometimes exceeded 1—which corresponds to complete interfacial polarization. The origin of this discrepancy remains to be fully resolved (Fert and Lee, 1996; Geux *et al.*, 2000; Hershfield and Zhao, 1997; Johnson, 1993b, 2002a). Results obtained from the Hanle effect, on similar samples, gave the expected $P_\Sigma < 1$ values (Johnson, 2002a).⁶²

A modification of the bipolar spin switch structure was used to demonstrate the spin injection into a niobium film (Johnson, 1994), realizing the theoretical assertion of Aronov (1976a) that nonequilibrium spin could be injected into a superconductor. Two insulating Al_2O_3 films were inserted between F1 and F2 (both made of Py) and a Nb film [see Fig. 5(b)]. The measurements were performed near the superconducting transition temperature

⁵⁸ Recall from the discussion leading to Eq. (39) that the spin-coupled signal $\propto \delta M$.

⁵⁹ The range of the angle ϕ , in the inset, is corrected from the one originally given in Fig. 3 of Johnson and Silsbee (1985).

⁶⁰ The fitting parameters are τ_s , P_Σ , and α (Johnson and Silsbee, 1988d), and since the diffusion coefficient is obtained from Einstein’s relation L_s is known.

⁶¹ $d \sim 100$ nm was much smaller than the separation between F1 and F2 in bulk Al wires (Johnson and Silsbee, 1985), and the amplitude of the Hanle effect was about 10^4 larger (Johnson, 2002a).

⁶² Theoretical estimates for V_s from which $P_\Sigma > 1$ was inferred are modified when one considers the Coulomb interaction and proximity effects—near the N/F interface the spin splitting of the carrier bands in the N region will be finite even at equilibrium. Model calculations (Chui, 1995; Chui and Cullen, 1995), which treat the F/N/F junction as a whole, show that the magnetic susceptibility χ in N can be much smaller than the free electron value and can increase the predicted $V_s \propto 1/\chi$. These corrections to the free-electron picture of an F/N/F junction are smaller for larger d , as in the bulk-wire geometry of Johnson and Silsbee (1985), where theoretical estimates of V_s did not lead to $P_\Sigma > 1$.

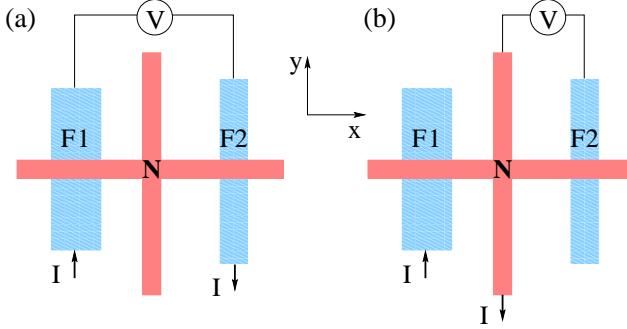


FIG. 11 Schematic representation of (a) local and (b) nonlocal geometry used to measure the effects of spin injection and spin accumulation.

T_c with the data qualitatively similar, above and below T_c , to the spin-coupled voltage, as obtained in the magnetic-field sweep from Fig. 10. The results were interpreted as support for enhanced depletion of the superconducting condensate (and correspondingly the reduction of the critical current I_c) by spin-polarized quasiparticles, as compared to the usual spin-unpolarized quasiparticle injection. Related measurements were recently performed in a CPP geometry (Gu *et al.*, 2002), and the penetration depth of the quasiparticle in the Nb films was measured to be ~ 16 nm, as compared to 2 nm in (Johnson, 1994). The corresponding temperature dependence of CPP GMR is well explained by the theory of Yamashita *et al.* (2003a) and the modification of Andreev reflection (see Sec. IV.A.3) by spin polarization.

The spin injection technique of Johnson and Silsbee was also applied to semiconductors. Initial experiments on using a metallic ferromagnet to inject spin into a two-dimensional electron gas (2DEG) showed only a very low ($\sim 1\%$) efficiency (Hammar *et al.*, 1999) for which various explanations were offered (Hammar *et al.*, 2000; Monzon *et al.*, 2000; van Wees, 2000). However, stimulated by the proposal of Rashba (2000) to employ spin-selective diffusive contacts (Sec. II.C.1), the subsequent measurements have showed substantially more efficient spin injection into a 2DEG after an insulating layer was inserted (Hammar and Johnson, 2001, 2002). The geometry employed is depicted in Fig. 9. In interpreting the results, the spin-orbit coupling and the energy-independent density of states at the Fermi level were taken into account (Silsbee, 2001). This topic is reviewed by Tang *et al.* (2002).

2. Spin injection into metals

An important part of the operation of CPP GMR structures is the presence of nonequilibrium spin polarization in nonmagnetic metallic regions. Studies of spin-injection parameters in such systems have been reviewed by Bass and Pratt, Jr. (1999) and Gijs and Bauer (1997). However, until recently, except for the work of Johnson

and Silsbee, there were few other experimental studies directly concerned with spin injection into metals. A series of experiments, (Jedema *et al.*, 2002a, 2001, 2002b,d) at both low (4.2 K) and room temperature, were performed using the van der Pauw geometry depicted in Fig. 11. In various structures (Jedema, 2002) the two ferromagnetic regions (made of Py, Co, or Ni) were chosen to be of different sizes to provide different coercive fields, allowing an independent reversal of magnetization in F1 and F2. The cross-shaped nonmagnetic region was made of Al or Cu (Jedema, 2002). Nonlocal measurements, similar to the approach shown in Figs. 5 and 9 [discussed in (Johnson, 1993b; Johnson and Silsbee, 1988d)], were shown to simplify the extraction of spurious effects (for example, anisotropic magnetoresistance and the Hall signal) from effects intrinsic to spin injection, as compared to the local or conventional spin-valve geometry.

In the first type of experiment the cross-shaped region was deposited directly over the F region (Fig. 11), and the spin-coupled resistance ΔR , defined analogously to Eq. (1), was measured as a function of an in-plane magnetic field. A theoretical analysis (Jedema *et al.*, 2001, 2002d) was performed assuming no interfacial resistance ($r_c = 0$) and the continuity of the electrochemical potentials at the F/N interface (see Sec. II.C.1). For a spin injection from Py into Cu, the maximum current polarization was obtained to be $P_j \approx 0.02$ at 4.2 K. The results for ΔR (Jedema *et al.*, 2001) scaled to the size of the samples used by Johnson (1993a,b) were interpreted to be 3-4 orders of magnitude smaller. As discussed in Secs. II.C.1 and II.C.2, the presence of interfacial spin-selective resistance can substantially change the spin injection efficiency and influence the resistance mismatch between the F and N regions [see Eq. (19)]. Estimates of how these considerations would affect the results of Jedema *et al.* (2001) were given by Jedema *et al.* (2002c) as well as by others (Johnson and Byers, 2003; Takahashi and Maekawa, 2003), who analyzed the importance of multidimensional geometry. In addition to comparing characteristic values of contact resistance obtained on different samples,⁶³ for a conclusive understanding it will be crucial to have *in situ* measurements.

In analyzing data for the van der Pauw cross, a two-dimensional geometry has an important effect—while the electric current is following the paths depicted in Fig. 11, the spin current, through the diffusion of nonequilibrium spin, would have similar flow in all four arms (Johnson, 2002a). This is different from the usual (quasi)one-dimensional analysis in which spin and charge currents flow along the same paths. For a full understanding of the van der Pauw cross geometry, two-dimensional modeling might be necessary (Johnson and Byers, 2003; Takahashi

⁶³ For example, the measured resistance of clean F/N contacts in CPP GMR (Bussmann *et al.*, 1998) was used to infer that there is also a large contact resistance in all-metal spin injection experiments (Johnson, 2002b).

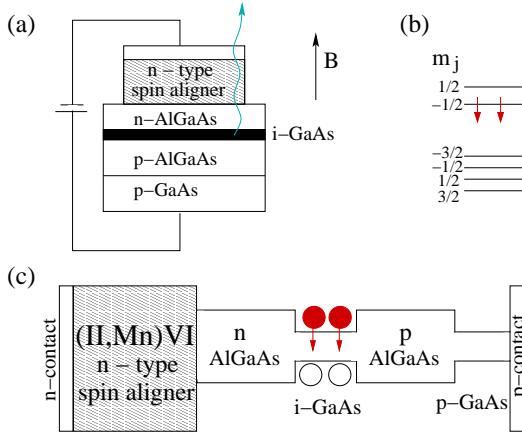


FIG. 12 Schematic device geometry and band diagram of a spin LED: (a) Recombination of spin-polarized electrons injected from the (II,Mn)VI spin aligner and unpolarized holes injected from the p-doped GaAs, in the intrinsic GaAs quantum well, producing circularly polarized light; (b) conduction and valence bands of a spin aligner in an external magnetic field; (c) sketch of the corresponding band edges and band offsets in the device geometry. In the quantum well, spin down electrons and unpolarized holes are depicted by solid and empty circles, respectively. Adapted from Fiederling *et al.*, 1999.

and Maekawa, 2003).

In the second type of experiment, tunneling contacts were fabricated by inserting Al_2O_3 as an insulator into the regions where F1 and F2 overlapped with the cross. By applying a transverse field B_z (see Fig. 11) the precession of the injected nonequilibrium spin was controlled and the amplitude of the Hanle effect was measured (Jedema *et al.*, 2002a,b), as outlined in Sec. II.D.1. From $\text{Co}/\text{Al}_2\text{O}_3/\text{Al}/\text{Al}_2\text{O}_3/\text{Co}$ structures $L_s \approx 0.5 \mu\text{m}$ was extracted at room temperature. The analysis of the Hanle signal was performed by averaging contributions of different lifetimes (D'yakonov and Perel', 1984b). This proved to be equivalent to the Johnson and Silsbee (1988d) solution to the Bloch-Torrey equations.

3. All-semiconductor spin injection

If a magnetic semiconductor could be used as a robust spin injector (spin aligner) into a nonmagnetic semiconductor it would facilitate the integration of spintronics and semiconductor-based electronics. Comparable resistivities of magnetic and nonmagnetic semiconductors could provide efficient spin injection [see Eq. (31), with $r_F \approx r_N$] even without using resistive contacts. Ultimately, for a wide range of applications and for compatibility with complementary metal-oxide semiconductors (CMOS) (Wong *et al.*, 1999), it would be desirable to be able to inject spin into silicon at room temperature.

Early studies (Osipov *et al.*, 1990, 1998; Viglin *et al.*, 1991, 1997), which have since largely been ignored, used a Cr- and Eu-based chalcogenide ferromagnetic semi-

conductor (FSm) (Nagaev, 1983) as the spin injector.⁶⁴ The experiments were motivated by the theoretical work of Aronov (1976a); Aronov and Pikus (1976) predicting that the ESR signal, proportional to the steady-state magnetization, would be changed by spin injection. The measurements of Osipov *et al.* (1990); Viglin *et al.* (1991) prompted a related prediction (Margulis and Margulis, 1994) that spin injection could be detected through changes in electric dipole spin resonance (EDSR). EDSR is the spin-flip resonance absorption for conduction electrons at Zeeman frequency, which is excited by the electric-field vector of an incident electromagnetic wave. The theory of EDSR, developed by Rashba and Sheka (1961) is extensively reviewed by Rashba and Sheka (1991).

Ferromagnetic semiconductor spin injectors formed $p-n$ and $n-n$ heterostructures with a nonmagnetic semiconductor InSb. The choice of InSb is very suitable due to its large negative (~ -50) g factor (McCombe and Wagner, 1971), for detecting the effects of spin injection through ESR. The observed absorption and emission of microwave power (Osipov *et al.*, 1998) was tuned by an applied magnetic field (from 35 GHz at ≈ 400 G up to 1.4 THz at 20 kG) and only seen when electrons flowed from FSm into an Sm region. The injection to the lower Zeeman level increased the ESR absorption, while injection to the higher Zeeman level, leading to population inversion.

The most recent experiments using semiconductor spin injectors can be grouped into two different classes, In one approach (II,Mn)VI paramagnetic semiconductors were employed as the spin aligners. These included CdMnTe (Oestreich *et al.*, 1999), BeMnZnSe (Fiederling *et al.*, 1999), and ZnMnSe (Jonker *et al.*, 2000). In the second approach ferromagnetic semiconductors like (Ga,Mn)As (Chun *et al.*, 2002; Mattana *et al.*, 2003; Ohno *et al.*, 1999b) were used. Both approaches were also employed to inject spins into CdSe/ZnSe (Seufert *et al.*, 2004) and InAs (Chye *et al.*, 2002) quantum dots, respectively.

In (II,Mn)VI materials, at low Mn-concentration and at low temperatures, there is a giant Zeeman splitting $\Delta E = g^* \mu_B H$ (Furdyna, 1988; Gaj, 1988) of the conduction band, in which g^* is the effective electron g factor. Such splitting arises due to $sp-d$ exchange between the spins of conduction electrons and the $S=5/2$ spins of the localized Mn^{2+} ions. The g^* factor for $H \neq 0$ can exceed⁶⁵ 100 and is given by (Brandt and Moshchalkov,

⁶⁴ These materials, while more difficult to fabricate than the subsequent class of III-V ferromagnetic semiconductors, have desirable properties of providing injection of spin-polarized electrons (with spin lifetimes typically much longer than for holes) and large spin splitting [~ 0.5 eV at 4.2 K for n-doped HgCr_2Se_4 (Nagaev, 1983)] with nearly complete spin polarization and a Curie temperature T_C of up to 130 K (HgCr_2Se_4) (Osipov *et al.*, 1998).

⁶⁵ At low temperatures (~ 1 K) $\text{Cd}_{0.95}\text{Mn}_{0.05}\text{Se}$ has $|g^*| > 500$

1984; Furdyna, 1988)

$$g^* = g + \alpha M / (g_{Mn} \mu_B^2 H), \quad (50)$$

where g is the $H = 0$ II-VI “band” value g , generally different from the free-electron value, magnetization $M \propto \langle S_z \rangle \propto B_s [(g_{Mn} \mu_B S H) / (k_B T)]$, B_s is the Brillouin function (Ashcroft and Mermin, 1976), and α is the exchange integral for s -like Γ_6 electrons (see Table I in Sec. II.B), given by (Furdyna, 1988)

$$\alpha \equiv \langle S | J_{sp-d} | S \rangle / V_0, \quad (51)$$

where J_{sp-d} is the electron-ion exchange coupling, and V_0 is the volume of an elementary cell. From Eqs. (50) and (51) it follows that $g^* = g^*(H)$ can even change its sign. Similar analysis applies also to g factors of holes, with the Zeeman splitting of a valence band being typically several times larger than that of a conduction band (Brandt and Moshchalkov, 1984).

(II,Mn)VI materials can be incorporated in high quality heterostructures with different optically active III-V nonmagnetic semiconductors which, by providing circularly polarized luminescence, can also serve as spin detectors. In this case carriers are excited by electrical means and we speak of electroluminescence rather than photoluminescence. The selection rules for the recombination light are the same as discussed in Sec. II.B.

Figure 12 depicts a scheme for realization of all-semiconductor electrical spin injection and optical detection (Fiederling *et al.*, 1999; Jonker *et al.*, 2000). Displayed is a spin light-emitting diode (LED) (Jonker, 1999) in a Faraday geometry where both the applied B -field and the direction of propagation of the emitted light lie along the growth direction. Similar to an ordinary LED (Sze, 1981), electrons and holes recombine (in a quantum well or a $p-n$ junction) and produce electroluminescence. However, in a spin LED, as a consequence of radiative recombination of spin-polarized carriers, the emitted light is circularly polarized. In experiments of Fiederling *et al.* (1999); Jonker *et al.* (2000), at $B \approx 1$ T, $T \approx 4$ K, and forward bias, electrons entering from the n -contact were almost completely polarized in the spin down state as they left the spin aligner and are injected across the (II,Mn)VI/AlGaAs interface. The electrons further traveled (by drift and diffusion) to an intrinsic GaAs quantum well (QW) where they recombined with the unpolarized holes, which were injected from the p -doped GaAs.⁶⁶

The efficiency of electrical spin injection across the (II,Mn)VI/AlGaAs interface was studied (Fiederling *et al.*, 1999) using P_{circ} (defined in Sec. II.B) of electroluminescence, as a function of B and the thickness of the magnetic spin aligner (0 nm, 3 nm, and 300 nm, respectively). P_{circ} increased with the thickness of the magnetic layer, suggesting the finite spin relaxation time needed for initially unpolarized electrons to relax into the lower (spin down) Zeeman level. The results of Jonker *et al.* (2000) were similar to those of Fiederling *et al.* (1999) for the thickest magnetic region. The behavior of $P_{\text{circ}}(B)$, up to the saturation value ($B \approx 3$ T), could be well explained by the magnetization described with the Brillouin function (Furdyna, 1988; Gaj, 1988), expected for the (II,Mn)VI semiconductors. In Fig. 12 the injected spin down electrons are majority electrons with their magnetic moments parallel to the applied magnetic field. The principles of optical orientation discussed in Sec. II.B and the selection rules for GaAs sketched in Fig. 6 are used to infer P_n in a QW.

For a QW of approximately the same width (150 nm) the conversion of P_{circ} to P_n used by Fiederling *et al.* (1999) differed by a factor of 2 from that used by Jonker *et al.* (2000). Fiederling *et al.* (1999) assumed that confinement effects were negligible, leading to the selection rules for a bulk GaAs (recall $P_{\text{circ}} = -P_n/2$, from Sec. II.B). The maximum $P_{\text{circ}} \approx 43\%$ was interpreted as implying nearly 90% polarized injected electrons. Jonker *et al.* (2000) inferred $|P_n| \approx 50\%$, from $P_{\text{circ}} = -P_n$ (Weisbuch and Vinter, 1991), as a consequence of QW confinement and lifting of the degeneracy between light and heavy hole states in the valence band ($\approx 5 - 6$ meV), see Fig. 6. Both results clearly demonstrated a robust low-temperature spin injection using the spin LED’s. Subsequent studies (Jonker *et al.*, 2001; Park *et al.*, 2000; Stroud *et al.*, 2002) have supported the lifting of degeneracy between the light and heavy hole bands. The corresponding data are shown in Fig. 13. Similar values of P_{circ} were also measured in a resonant tunneling diode based on ZnMnSe (Gruber *et al.*, 2001; Waag *et al.*, 2001). Spin injection using the spin LED’s, described above, is not limited to structures grown by molecular-beam epitaxy (MBE). It is also feasible using air-exposed interfaces (Park *et al.*, 2000) similar to the actual fabrication conditions employed in conventional electronics.

The robustness of measured P_{circ} was studied by intentionally changing the density of linear defects, from stacking faults at the ZnMnSe/AlGaAs interface (Stroud *et al.*, 2002). An approximate linear decrease of P_{circ} with the density of stacking faults was shown to be consistent with the influence of spin-orbit interaction as modeled by Elliot-Yafet scattering (see Sec. III.B.1) at the interface. The nonspherically symmetric defect potential (entering the spin-orbit interaction) causes a highly anisotropic loss of spin polarization. At small angles to the axis of growth [see Fig. (12)], the probability of the spin flip of an injected electron is very high, leading to a small spin po-

(Dietl, 1994), while in n-doped (In,Mn)As $|g^*| > 100$ at 30 K (Zudov *et al.*, 2002). Such large g factors, in the presence of a highly inhomogeneous magnetic field could lead to the charge carrier localization (Berciu and Jankó, 2003).

⁶⁶ The spatial separation and spin relaxation between the spin injection and the point of spin detection (in QW) make a fully quantitative analysis of the injected polarization more difficult. It would be valuable to perform realistic calculations of a spin-polarized transport and spin injection which would treat the whole spin LED as a single entity.

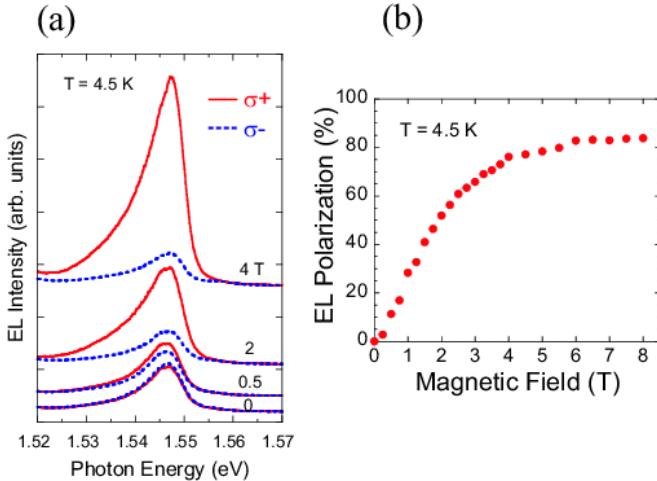


FIG. 13 (a) Electroluminescence in a spin light-emitting diode (LED): (a) Electroluminescence (EL) spectra from a surface-emitting spin LED with a $Zn_{0.94}Mn_{0.06}Se$ contact for selected values of applied magnetic field, analyzed for σ^\pm (positive and negative helicity); the magnetic field is applied along the surface normal (Faraday geometry) and the spectra are dominated by the heavy hole exciton; (b) magnetic field dependence of the EL circular polarization. Adapted from Jonker *et al.*, 2001.

larization. These findings illustrate the importance of interface quality and the effect of defects on the spin injection efficiency, an issue not limited to semiconductor heterostructures. Related information is currently being sought by spatial imaging of the spin polarization in spin LED's (Thruber and Smith, 2003; Thruber *et al.*, 2002) using magnetic resonance force microscopy (Sidles *et al.*, 1995).

(III,Mn)V ferromagnetic semiconductors are also used to inject spin in a spin-LED structures as depicted in Fig. 12. Spin injection can be achieved with no external field, and the reports of high T_C in some compounds suggest that all-semiconductor spin LED's could operate at room temperature. The drawback, however, is that the most (III,Mn)V's have spin-polarized holes (rather than electrons) as the main carriers which, due to spin-orbit coupling, lose their polarization very quickly after being injected into a nonmagnetic semiconductor. Consequently, results of the spin injection show only a small degree of hole polarization.

In the experiment of Ohno *et al.* (1999b), an intrinsic GaAs spacer of thickness d was introduced between the spin aligner (Ga,Mn)As and the (In,Ga)As quantum well. The electroluminescence in a QW was measured perpendicular to the growth direction [the easy magnetization axis of (Ga,Mn)As and the applied magnetic field were both perpendicular to the growth direction]. The corresponding relation between the P_{circ} and hole density polarization P_p is not straightforward; the analysis was performed only on the electroluminescence [for possible difficulties see Fiederling *et al.* (2003)]. A small measured signal ($P_{\text{circ}} \sim 1\%$ at 5 K), consistent with the expecta-

tion for holes as the injected spin-polarized carriers, was also obtained in an additional experiment (Young *et al.*, 2002). P_{circ} was approximately independent of the GaAs thickness ($d = 20 - 420$ nm), a behavior which remains to be understood considering that the hole spins should relax fast (Hilton and Tang, 2002) as they are transferred across the nonmagnetic semiconductor.⁶⁷ In contrast, for a repeated experiment (Young *et al.*, 2002) using a Faraday geometry (as in Fig. 12), with both measured electroluminescence and B along the growth direction, the same change of thickness P_{circ} was reduced from 7% to 0.5%. A highly efficient spin injection of $P_n \approx 80\%$ in GaAs has been realized using (Ga,Mn)As as a spin injector in a Zener diode structure (Van Dorpe *et al.*, 2003a). The detection employed the technique of an oblique Hanle effect, discussed in the next section.

All-electrical spin injection studies of trilayer structures (II,Mn)VI/II-VI/(II,Mn)VI have displayed up to 25% MR at $B \approx 5$ T and $T = 4$ K (Schmidt *et al.*, 2001). A strong suppression of this MR signal at applied bias of ~ 10 mV was attributed to the nonlinear regime of spin injection, in which the effects of band bending and charge accumulation at the (II,Mn)VI/II-VI interface were important (Schmidt *et al.*, 2002). It would be instructive to analyze these measurements by adopting the approach discussed in the context of magnetic p - n junctions (Secs. II.C.3 and IV.D), which self-consistently incorporates the effects of band bending and deviation from local charge neutrality.

4. Metallic ferromagnet/semiconductor junctions

A large family of metallic ferromagnets, some of them highly spin polarized, offer the possibility of spin injection at room temperature, even in the absence of applied magnetic field. Spin injection into (110) GaAs at room temperature has been already demonstrated using vacuum tunneling from a polycrystalline Ni STM tip and optical detection via circularly polarized luminescence (Alvarado, 1995; Alvarado and Renaud, 1992). It was shown that the minority spin electrons (spin \downarrow in the context of metals; see Sec. II.A) in Ni produced the dominant contribution to the tunneling current, and the resulting polarization was inferred to be $P_n = (-31 \pm 5.6)\%$ (Alvarado and Renaud, 1992). Even though the spin injection in future spintronic devices will likely be implemented by some means other than vacuum tunneling, this result supports the importance of the tunneling contact for efficient spin injection, as discussed in Sec. II.C.1. Similar studies of spatially resolved spin injection, sensitive to the topography of the GaAs surface, have employed a single-crystal Ni (100) tip (LaBella *et al.*, 2001).

⁶⁷ A possible exception is QW, in which the effects of quantum confinement and quenching spin-orbit coupling lead to longer τ_s .

At 100 K nearly fully spin-polarized injection of electrons was reported. However, further analyses of the measurements of P_{circ} have substantially reduced these estimates to $\approx 25\%$ (Egelhoff, Jr. *et al.*, 2002; LaBella *et al.*, 2002).

Direct spin injection from a ferromagnet into a 2DEG,⁶⁸ motivated by the proposal of Datta and Das (1990) proposal, initially showed only small effects (Gardelis *et al.*, 1999; Hammar *et al.*, 1999; Lee *et al.*, 1999), with $\Delta R/R \sim 1\%$, or effects within the noise (Filip *et al.*, 2000). Such inefficiency could be attributed to the resistance mismatch in the F and N regions, discussed in Secs. II.C.1 and II.C.2. The possibility of spurious effects arising from the Hall and anisotropic magnetoresistance signals in similar structures was suggested earlier (Monzon *et al.*, 1997) as well as after the initial experiments (Monzon *et al.*, 2000; Tang *et al.*, 2002; van Wees, 2000). Control measurements have been performed to address these issues (Hammar *et al.*, 2000; Hammar and Johnson, 2000). This debate about the presence/absence of spin injection effects via Ohmic contacts stimulated further studies, but the experimental focus has since shifted to other approaches.

Spin injection via Schottky contacts at room temperature was demonstrated in a Fe/GaAs junction by Zhu *et al.* (2001), who reported detection of $P_{\text{circ}} \approx 2\%$ using spin LED structures and optical detection as described in II.D.3. These studies were extended (Ramsteiner *et al.*, 2002) by using MBE to grow MnAs, a ferromagnetic metal, on top of the GaAs to provide high-quality interfaces (Tanaka, 2002). There was no preferential behavior for spin injection using different azimuthal orientations of the epitaxial MnAs layer, which could have been expected from the symmetry between conduction-band wave functions in MnAs and GaAs. The tunneling properties of a Schottky barrier were discussed by Gibson and Meservey (1985); Kreuzer *et al.* (2002); Meservey *et al.* (1982); Prins *et al.* (1995). The measured I-V curves display a complicated behavior (Hirohata *et al.*, 2001; Isaković *et al.*, 2001) which can be significantly affected by the interface (midgap) states at the Schottky barrier (Jonker *et al.*, 1997). A theoretical explanation of this behavior is still lacking.

As discussed in Sec. II.C.1, tunnel contacts formed between a metallic ferromagnet and a semiconductor can provide effective spin injections. Optical detection in spin LED structures, as discussed in Sec. II.D.3, was used to show carrier polarization of $P_n \approx 30\%$ using Fe as a spin injector (Hanbicki *et al.*, 2003). Experimental results are given in Fig. 14. Some care has to be taken in defining the efficiency of the spin injection, normalized to the polarization of a ferromagnet, as used in related previous experiments (Hanbicki and Jonker, 2002; Hanbicki *et al.*, 2002; Jansen, 2002). Furthermore, there are often different conventions for defining the sign of P_n (Hanbicki

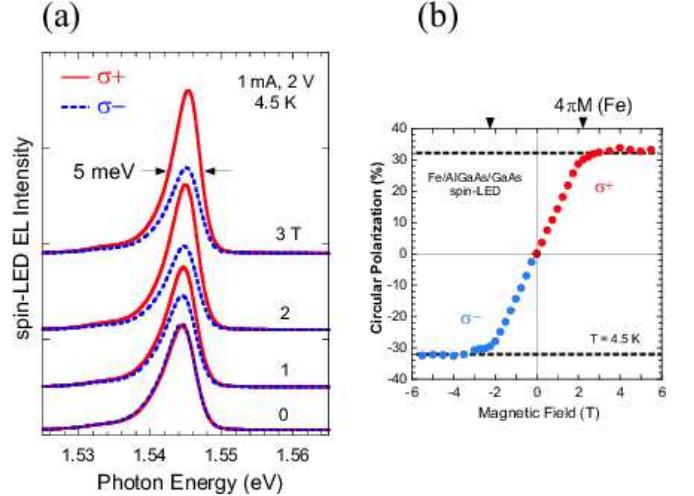


FIG. 14 Electroluminescence (EL) and polarization due to spin injection from Fe Schottky contact: (a) EL spectra from a surface-emitting spin LED with an Fe/AlGaAs Schottky tunnel contact for selected values of applied magnetic field, analyzed for σ^\pm circular polarization. The large difference in intensity between these components indicates successful spin injection from the Fe into the GaAs QW, and reveals an electron spin polarization in the QW of 32%. The magnetic field is applied along the surface normal (Faraday geometry). The spectra are dominated by the heavy hole (HH) exciton. Typical operating parameters are 1 mA and 2 V. (b) Magnetic-field dependence of the EL circular polarization of the HH exciton. The polarization tracks the hard axis magnetization of the Fe contact, and saturates at an applied magnetic-field value $4\pi M = 2.2$ T, at which the Fe magnetization is entirely along the surface normal. From Hanbicki *et al.*, 2003.

et al., 2003), used in the context of semiconductors and ferromagnetic metals, as discussed in Sec. II.A. Jiang *et al.* (2004) have demonstrated that MgO can be a suitable choice of an insulator for highly efficient spin injection into GaAs. Spin LED with GaAs/AlGaAs quantum well was used to detect $P_n \approx 50\%$ at 100 K injected from CoFe/MgO (100) tunnel injector. While quantum well emission efficiency limits detection at higher temperatures (> 100 K), the same tunnel injector should also be suitable for efficient spin injection even at room temperature.

An oblique Hanle effect (D'yakonov *et al.*, 1974) (see also Secs. II.B and II.D.1) was used (Motsnyi *et al.*, 2002, 2003; Van Dorpe *et al.*, 2003b) to detect spin injection, giving up to $P_n \approx 16\%$, at room temperature. The geometry used is similar to that sketched in Fig. 12(a), with an insulating layer (AlO_x) separating the ferromagnetic spin injector and the (Al,Ga)As/GaAs spin LED. The magnetization easy axis lies in the plane of the ferromagnet. An oblique magnetic field is applied to give a net out-of-plane component of injected spin which could contribute to the emission of circularly polarized light. This approach allows one to apply a magnetic field several times smaller than would be needed to pull the magnetization out of plane [for Fe it is ≈ 2 T (Hanbicki *et al.*, 2002)] and to

⁶⁸ For a comprehensive review of 2DEG, see Ando *et al.* (1982).

measure polarized luminescence in a Faraday geometry. Furthermore, using standard measurements of the Hanle curve, one can extract separately the spin lifetime and carrier recombination time.

Hot-electron spin injection above the Schottky barrier is another method for using a high polarization of metallic ferromagnets to create a nonequilibrium spin in a semiconductor even at room temperatures. Typically such injection is performed in three-terminal, transistor-like devices, as discussed in Sec. IV.E.3.

Direct electrical spin injection has also been demonstrated in organic semiconductors (Dedić *et al.*, 2002). MR measurements were performed in an F/N/F junction, where F is $\text{La}_{0.7}\text{Sr}_{0.3}\text{MnO}_3$ (LSMO), a colossal magnetoresistive manganite, and N is sexithienyl (T_6), a π -conjugated rigid-rod oligomer organic semiconductor. The decrease of MR with increasing thickness of the N region was used to infer $L_{SN} \sim 100$ nm at room temperature. The resulting spin diffusion length is a combination of low mobility, $\sim 10^{-4}\text{cm}^2\text{V}^{-1}\text{s}^{-1}$ (about $\sim 10^7$ times smaller than for the bulk GaAs), and long spin relaxation times, $\sim \mu\text{s}$,⁶⁹ as compared to the usual III-V inorganic semiconductors. Motivated by these findings Xiong *et al.* (2004) have replaced one of the LSMO electrodes by Co. Different coercive fields in the two ferromagnetic electrodes allowed them to measure a spin-valve effect with $\Delta R/R \sim 40\%$ at 11 K. Related theoretical studies of the ferromagnetic metal/conjugated polymer interfaces were reported by Xie *et al.* (2003).

III. SPIN RELAXATION AND SPIN DEPHASING

A. Introduction

Spin relaxation and spin dephasing are processes that lead to spin equilibration and are thus of great importance for spintronics. The fact that nonequilibrium electronic spin in metals and semiconductors lives relatively long (typically a nanosecond), allowing for spin-encoded information to travel macroscopic distances, is what makes spintronics a viable option for technology. After introducing the concepts of spin relaxation and spin dephasing times T_1 and T_2 , respectively, which are commonly called τ_s throughout this paper, we discuss four major physical mechanisms responsible for spin equilibration in *nonmagnetic* electronic systems: Elliott-Yafet, D'yakonov-Perel', Bir-Aronov-Pikus, and hyperfine-interaction processes. We then survey recent works on electronic spin relaxation in nonmagnetic metals and semiconductors, using the important examples of Al and GaAs for illustration.

1. T_1 and T_2

Spin relaxation and spin dephasing of a spin ensemble are traditionally defined within the framework of the Bloch-Torrey equations (Bloch, 1946; Torrey, 1956) for magnetization dynamics. For mobile electrons, spin relaxation time T_1 (often called longitudinal or spin-lattice time) and spin dephasing time T_2 (also transverse or decoherence time) are defined via the equations for the spin precession, decay, and diffusion of electronic magnetization \mathbf{M} in an applied magnetic field $\mathbf{B}(t) = B_0\hat{\mathbf{z}} + \mathbf{B}_1(t)$, with a static longitudinal component B_0 (conventionally in the $\hat{\mathbf{z}}$ direction) and, frequently, a transverse oscillating part \mathbf{B}_1 perpendicular to $\hat{\mathbf{z}}$ (Kaplan, 1959; Torrey, 1956):

$$\frac{\partial M_x}{\partial t} = \gamma(\mathbf{M} \times \mathbf{B})_x - \frac{M_x}{T_2} + D\nabla^2 M_x, \quad (52)$$

$$\frac{\partial M_y}{\partial t} = \gamma(\mathbf{M} \times \mathbf{B})_y - \frac{M_y}{T_2} + D\nabla^2 M_y, \quad (53)$$

$$\frac{\partial M_z}{\partial t} = \gamma(\mathbf{M} \times \mathbf{B})_z - \frac{M_z - M_z^0}{T_1} + D\nabla^2 M_z. \quad (54)$$

Here $\gamma = \mu_B g/\hbar$ is the electron gyromagnetic ratio (μ_B is the Bohr magneton and g is the electronic g factor), D is the diffusion coefficient (for simplicity we assume an isotropic or a cubic solid with scalar D), and $M_z^0 = \chi B_0$ is the thermal equilibrium magnetization with χ denoting the system's static susceptibility. The Bloch equations are phenomenological, describing quantitatively very well the dynamics of mobile electron spins (more properly, magnetization) in experiments such as conduction electron spin resonance and optical orientation. Although relaxation and decoherence processes in a many-spin system are generally too complex to be fully described by only two parameters, T_1 and T_2 are nevertheless an extremely robust and convenient measure for quantifying such processes in many cases of interest. To obtain microscopic expressions for spin relaxation and dephasing times, one starts with a microscopic description of the spin system (typically using the density-matrix approach), derives the magnetization dynamics, and compares it with the Bloch equations to extract T_1 and T_2 .

Time T_1 is the time it takes for the longitudinal magnetization to reach equilibrium. Equivalently, it is the time of thermal equilibration of the spin population with the lattice. In T_1 processes an energy has to be taken from the spin system, usually by phonons, to the lattice. Time T_2 is classically the time it takes for an ensemble of transverse electron spins, initially precessing in phase about the longitudinal field, to lose their phase due to spatial and temporal fluctuations of the precessing frequencies. For an ensemble of mobile electrons the measured T_1 and T_2 come about by averaging spin over the thermal distribution of electron momenta. Electrons in different momentum states have not only different spin-flip characteristics, but also slightly different g factors and thus different precession frequency. This

⁶⁹ This is also a typical value for other organic semiconductors (Krinichnyi, 2000), a consequence of weak spin-orbit coupling (Davis and Bussmann, 2003).

is analogous to precession frequencies fluctuations of localized spins due to inhomogeneities in the static field B_0 . However, since momentum scattering (analogous to intersite hopping or exchange interaction of localized spins) typically proceeds much faster than spin-flip scattering, the g -factor-induced broadening is inhibited by motional narrowing⁷⁰ and need not be generally considered as contributing to T_2 [see, however, (Dupree and Holland, 1967)]. Indeed, motional narrowing of the g -factor fluctuations, δg , gives a contribution to $1/T_2$ of the order of $\Delta\omega^2\tau_p$, where the B_0 -dependent precession frequency spread is $\Delta\omega = (\delta g/g)\gamma B_0$ and τ_p is the momentum scattering time. For B_0 fields of the order 1 T, scattering times of 1 ps, and δg as large as 0.01, the “inhomogeneous broadening” is a microsecond, which is much more than the observed values for T_2 . Spatial inhomogeneities of B_0 , like those coming from hyperfine fields, are inhibited by motional narrowing, too, due to the itinerant nature of electrons. For localized electrons (e.g., for donor states in semiconductors), spatial inhomogeneities play an important role and are often observed to affect T_2 . To describe such reversible phase losses, which can potentially be eliminated by spin-echo experiments, sometimes the symbol T_2^* (Hu *et al.*, 2001b) is used to describe spin dephasing of ensemble spins, while the symbol T_2 is reserved for irreversible loss of the ensemble spin phase. In general, $T_2^* \leq T_2$, although for conduction electrons to a very good approximation $T_2^* = T_2$.

In isotropic and cubic solids $T_1 = T_2$ if $\gamma B_0 \ll 1/\tau_c$, where τ_c is the so-called correlation or interaction time: $1/\tau_c$ is the rate of change of the effective dephasing magnetic field (see footnote 70). Phase losses occur during time intervals of τ_c . As shown below, in electronic systems τ_c is given either by τ_p or by the time of the interaction of electrons with phonons and holes. Those times are typically smaller than a picosecond, so $T_1 = T_2$ is fulfilled for magnetic fields up to several tesla. The equality between the relaxation and dephasing times was noticed first in the context of NMR (Bloch, 1946; Wangsness and Bloch, 1953) and later extended to electronic spin systems (Andreev and Gerasimenko, 1958; Pines and Slichter, 1955). A qualitative reason for $T_1 = T_2$ is that if the phase acquires a random contribution during a short time interval τ_c , the energy uncertainty of the spin levels determining the longitudinal spin is greater than the

Zeeman splitting $\hbar\gamma B_0$ of the levels. The splitting then does not play a role in dephasing, and the dephasing field will act equally on the longitudinal and transverse spin. In classical terms, spin that is oriented along the direction of the magnetic field can precess a full period about the perpendicular fluctuating field, feeling the same dephasing fields as transverse components. As the external field increases, the precession angle of the longitudinal component is reduced, inhibiting dephasing.

The equality of the two times is very convenient for comparing experiment and theory, since measurements usually yield T_2 , while theoretically it is often more convenient to calculate T_1 . In many cases a single symbol τ_s is used for spin relaxation and dephasing (and called indiscriminately either of these terms), if it does not matter what experimental situation is involved, or if one is working at small magnetic fields.⁷¹ Throughout this paper we adopt this notation to avoid unnecessary confusion.

If the system is anisotropic, the equality $T_1 = T_2$ no longer holds, even in the case of full motional narrowing of the spin-spin interactions and g -factor broadening. Using simple qualitative analysis Yafet (1963) showed that, while there is no general relation between the two times, the inequality $T_2 \leq 2T_1$ holds, and that T_2 changes with the direction by at most a factor of 2. In the motionally narrowed case this difference between T_1 and T_2 can be considered as arising from the tensorial nature of the spin relaxation rate. Specific examples of this will be discussed in studying spin relaxation in two-dimensional semiconductor heterostructures (Sec. III.B.2).

Finally, we discuss the connection between τ_s and the single-spin decoherence time⁷², τ_{sc} , which is the single-spin correlation time. Time τ_{sc} becomes important for applications in spin-based quantum computing (Hu and Das Sarma, 2000; Loss and DiVincenzo, 1998), where spin coherence needs to last for at least $10^4 - 10^6$ gate operations for the computation to be fault tolerant (Preskill, 1998). The relative magnitudes of τ_s and τ_{sc} depend on many factors. Often $\tau_{sc} < \tau_s$, as is the case for a direct exchange interaction causing single-spin decoherence, while contributing to ensemble dephasing only as a dynamical averaging factor (the exchange interaction preserves total spin). An analogy with momentum scattering may be helpful. Electron-electron collisions lead to individual momentum equilibration while conserving the total momentum and hence do not contribute to charge current (unless umklapp processes are taken into account). A momentum scattering time obtained from conductivity (analogous to τ_s) would thus be very

⁷⁰ Motional (dynamical) narrowing is an inhibition of phase change by random fluctuations (Slichter, 1989). Consider a spin rotating with frequency ω_0 . The spin phase changes by $\Delta\phi = \omega_0 t$ over time t . If the spin is subject to a random force that makes spin precession equally likely clockwise and anticlockwise, the average spin phase does not change, but the root-mean-square phase change increases with time as $(\langle \Delta^2 \phi \rangle)^{1/2} \approx (\omega_0 \tau_c)(t/\tau_c)^{1/2}$, where τ_c is the correlation time of the random force, or the average time of spin precession in one direction. This is valid for rapid fluctuations, $\omega_0 \tau_c \ll 1$. The phase relaxation time t_ϕ is defined as the time over which the phase fluctuations reach unity: $1/t_\phi = \omega_0^2 \tau_c$.

⁷¹ Sometimes one finds as spin relaxation time $2\tau_s$. While this is correct for just one spin state, conventionally by spin relaxation we mean magnetization relaxation, in which each spin flip adds to the equilibration of both spin up and spin down states, doubling the rate for magnetization relaxation.

⁷² To distinguish ensemble and individual spin dephasing, we use the term decoherence in relation to single, or a few, spins.

different from a single-state momentum time (analogous to τ_{sc}). It is not clear at present how much τ_s and τ_{sc} differ for different materials under different conditions, although it is often, with little justification, assumed that they are similar. As Dzhioev *et al.* (2002c) recently suggested, τ_{sc} can be smaller than τ_s by several orders of magnitude in n-GaAs at low doping densities where electrons are localized in donor states, see also Sec. III.D.3.a. We note that it is τ_s that is relevant for spintronics (spin transport) applications, while τ_{sc} is relevant for solid-state quantum computing. Much remains to be learned about τ_{sc} .

2. Experimental probes

Experiments detecting spin relaxation and decoherence of conduction electrons can be grouped into two broad categories: (a) those measuring spectral characteristics of magnetization depolarization and (b) those measuring time or space correlations of magnetization.

Experiments of type (a) are exemplified by conduction-electron spin resonance (CESR) and optical orientation in combination with the Hanle effect. CESR was the first technique used to detect the spin of conduction electrons in metals (Feher and Kip, 1955; Griswold *et al.*, 1952) and donor states in semiconductors like Si (Feher and Gere, 1959). In GaAs, spin resonance techniques are aided by other measurements, e.g., optical detection (Weisbuch and Hermann, 1977), photoconductivity (Stein *et al.*, 1983), or magnetoresistance (Dobers *et al.*, 1988). The technique measures signatures of resonant absorption of microwaves by a Zeeman-split spin system of conduction electrons. Typically changes in surface impedance and in the transmission coefficient of the sample are observed. By comparing the absorption resonance curve with theory (Dyson, 1955; Kaplan, 1959) one can obtain both the carrier g factor and T_2 .

Optical spin orientation (Meier and Zakharchenya (Eds.), 1984) is a method of exciting spin-polarized carriers (electrons and holes) in direct-gap semiconductors like GaAs by absorption of circularly polarized light (see Sec. II.B). The injected spin polarization can be detected by observing circularly polarized luminescence resulting from recombination of the spin-polarized electrons and holes. Since in a steady state the excited spin polarization depends not only on τ_s but also on the carrier recombination time, the Hanle effect, polarization of luminescence by a transverse magnetic field (see Sec. II.D.1), is employed to deduce τ_s unambiguously.

The Hanle effect has also been a great tool for investigating T_2 in metals, in connection with electrical spin injection. The advantage of optical orientation over CESR is that, if carrier lifetime is known, zero-field (or zero- g -factor material) data can be measured. In addition, smaller τ_s values can be detected.

Type (b) experiments measure magnetization in a time or space domain. The most important examples are

the Johnson-Silsbee spin injection experiment, the time-resolved (pump-probe) photoluminescence (in which the probe is used on the same principle as optical orientation), and time-resolved Faraday and (magneto-optic) Kerr rotation. The last two methods can follow coherent (in the ensemble sense) dynamics of electron spin precession.

Spin injection experiments (Johnson and Silsbee, 1985) detect the amount of nonequilibrium magnetization by observing charge response (see Sec. II.D.1). In the Johnson-Silsbee scheme, electrons are first injected by electrical spin injection from a ferromagnetic electrode into a metal or semiconductor. As the spin diffuses throughout the sample, another ferromagnetic electrode detects the amount of spin as a position-dependent quantity. The detection is by means of spin-charge coupling, whereby an EMF appears across the paramagnet/ferromagnet interface in proportion to the nonequilibrium magnetization (Silsbee, 1980). By fitting the spatial dependence of magnetization to the exponential decay formula, one can extract the spin diffusion length L_s and thus the spin relaxation time $T_1 = L_s^2/D$. The Hanle effect, too, can be used in combination with Johnson-Silsbee spin injection, yielding directly $T_1 = T_2$, which agrees with T_1 determined from the measurement of L_s . A precursor to the Hanle effect in spin injection was transmission-electron spin resonance (TESR), in which nonequilibrium electron spin excited in the skin layer of a metallic sample is transported to the other side, emitting microwave radiation. In very clean samples and at low temperatures, electrons ballistically propagating through the sample experience Larmor precession resulting in Larmor waves, seen as an oscillation of the transmitted radiation amplitude with changing static magnetic field (Jánossy, 1980).

Time-resolved photoluminescence detects, after creation of spin-polarized carriers, circular polarization of the recombination light. This technique was used, for example, to detect a 500 ps spin coherence time T_2 of free excitons in a GaAs quantum well (Heberle *et al.*, 1994). The Faraday and (magneto-optic) Kerr effects are the rotation of the polarization plane of a linearly polarized light, transmitted through (Faraday) or reflected by (Kerr) a magnetized sample. The Kerr effect is more useful for thicker and nontransparent samples or for thin films fabricated on thick substrates. The angle of rotation is proportional to the amount of magnetization in the direction of the incident light. Pump (a circularly polarized laser pulse) and probe experiments employing magneto-optic effects can now follow, with 100 fs resolution, the evolution of magnetization as it dephases in a transverse magnetic field. Using lasers for spin excitation has the great advantage of not only detecting, but also manipulating spin dephasing, as shown using Faraday rotation on bulk GaAs and GaAs/ZnSe heterostructures (Awschalom and Kikkawa, 1999; Kikkawa *et al.*, 2001). The Kerr effect was used, for example, to investigate the spin dynamics of bulk CdTe (Kimel *et al.*, 2000), and

Faraday rotation was used to study spin coherence in nanocrystals of CdSe (Gupta *et al.*, 2002) and coherent control of spin dynamics of excitons in GaAs quantum wells (Heberle *et al.*, 1996).

B. Mechanisms of spin relaxation

Four mechanisms for spin relaxation of conduction electrons have been found relevant for metals and semiconductors: the Elliott-Yafet (EY), D'yakonov-Perel' (DP), Bir-Aronov-Pikus (BIP), and hyperfine-interaction (HFI) mechanisms.⁷³ In the EY mechanism electron spins relax because the electron wave functions normally associated with a given spin have an admixture of the opposite spin states, due to spin-orbit coupling induced by ions. The DP mechanism explains spin dephasing in solids without a center of symmetry. Spin dephasing occurs because electrons feel an effective magnetic field, resulting from the lack of inversion symmetry and from the spin-orbit interaction, which changes in random directions every time the electron scatters to a different momentum states. The BIP mechanism is important for p-doped semiconductors, in which the electron-hole exchange interaction gives rise to fluctuating local magnetic fields flipping electron spins. Finally, in semiconductor heterostructures (quantum wells and quantum dots) based on semiconductors with a nuclear magnetic moment, it is the hyperfine interaction of the electron spins and nuclear moments which dominates spin dephasing of localized or confined electron spins. An informal review of spin relaxation of conduction electrons can be found in Fabian and Das Sarma (1999c).

1. Elliott-Yafet mechanism

Elliott (1954) was the first to realize that conduction-electron spins can relax via ordinary momentum scattering (such as by phonons or impurities) if the lattice ions induce spin-orbit coupling in the electron wave function. In the presence of the spin-orbit interaction

$$V_{so} = \frac{\hbar}{4m^2c^2} \nabla V_{sc} \times \hat{\mathbf{p}} \cdot \hat{\boldsymbol{\sigma}}, \quad (55)$$

where m is the free-electron mass, V_{sc} is the scalar (spin-independent) periodic lattice potential, $\hat{\mathbf{p}} \equiv -i\hbar\nabla$ is the linear momentum operator, and $\hat{\boldsymbol{\sigma}}$ are the Pauli matrices, single-electron (Bloch) wave functions in a solid are no longer the eigenstates of $\hat{\sigma}_z$, but rather a mixture of the Pauli spin up $| \uparrow \rangle$ and spin down $| \downarrow \rangle$ states. If the solid possesses a center of symmetry, the case of elemental metals which Elliott considered, the Bloch states of

“spin up” and “spin down” electrons with the lattice momentum \mathbf{k} and band index n can be written as (Elliott, 1954)

$$\Psi_{\mathbf{k}n\uparrow}(\mathbf{r}) = [a_{\mathbf{k}n}(\mathbf{r})| \uparrow \rangle + b_{\mathbf{k}n}(\mathbf{r})| \downarrow \rangle] e^{i\mathbf{k}\cdot\mathbf{r}}, \quad (56)$$

$$\Psi_{\mathbf{k}n\downarrow}(\mathbf{r}) = [a_{-\mathbf{k}n}^*(\mathbf{r})| \downarrow \rangle - b_{-\mathbf{k}n}^*(\mathbf{r})| \uparrow \rangle] e^{i\mathbf{k}\cdot\mathbf{r}}, \quad (57)$$

where we write the explicit dependence of the complex lattice-periodic coefficients a and b on the radius vector \mathbf{r} . The two Bloch states are degenerate: they are connected by spatial inversion and time reversal (Elliott, 1954). That it makes sense to call $\Psi_{\mathbf{k}n\uparrow}$ and $\Psi_{\mathbf{k}n\downarrow}$, respectively, spin up and spin down states follows from the fact that in most cases the typical values of $|a|$ are close to unity, while $|b| \ll 1$.

Indeed, consider a band structure in the absence of spin-orbit coupling. Turning V_{so} on couples electron states of opposite Pauli spins which are of the same \mathbf{k} (because V_{so} has the period of the lattice), but different n . Because the spin-orbit interaction is normally much smaller than the distance between the bands, perturbation theory gives

$$|b| \approx \lambda_{so}/\Delta E \ll 1, \quad (58)$$

where ΔE is the energy distance between the band state in question and the state (of the same momentum) in the nearest band, and λ_{so} is the amplitude of the matrix element of V_{so} between the two states. The spin-orbit coupling alone does not lead to spin relaxation. However, in combination with momentum scattering, the spin up [Eq. (56)] and spin down [Eq. (57)] states can couple and lead to spin relaxation.

Momentum scattering is typically caused by impurities (at low T) and phonons (at high T). There is another important spin-flip scattering mechanism that involves phonons. A periodic, lattice ion-induced spin-orbit interaction is modified by phonons and can directly couple the (Pauli) spin up and spin down states. Such processes were first considered for a jellium model by Overhauser (1953a) [see also (Grimaldi and Fulde, 1997)], and for band structure systems by Yafet (1963). They must be combined with the Elliott processes discussed above to form a consistent picture of phonon-induced spin relaxation, especially at low temperatures (Yafet, 1963), where the two processes have similar magnitudes. At larger T , phonon-modified V_{so} is not important for polyvalent metals, whose spin relaxation is dominated by spin hot spots—states with anomalously large $|b|$ —as shown by the explicit calculation of Fabian and Das Sarma (1999b). Spin hot spots are discussed in more detail in Sec. III.C.

We now give a recipe, useful for elemental metals and semiconductors, for calculating phonon-induced $1/\tau_s$ from the known band and phonon structure. The corresponding theory was systematically developed by Yafet (1963). The spin relaxation rate due to phonon scattering according to the EY mechanism can be expressed through the spin-flip Eliashberg function $\alpha_S^2 F(\Omega)$ as (Fabian and

⁷³ We do not consider magnetic scattering, that is, scattering due to an exchange interaction between conduction electrons and magnetic impurities.

Das Sarma, 1999b)

$$1/\tau_s = 8\pi T \int_0^\infty d\Omega \alpha_s^2 F(\Omega) \partial N(\Omega)/\partial T, \quad (59)$$

where $N(\Omega) = [\exp(\hbar\Omega/k_B T) - 1]^{-1}$ is the phonon distribution function. The spin-flip Eliashberg function gives the contribution of the phonons with frequency Ω to the spin-flip electron-phonon interaction,

$$\alpha_s^2 F(\Omega) = \frac{g_S}{2M\Omega} \sum_\nu \langle \langle g_{kn\uparrow, k'n'\downarrow}^\nu \delta(\omega_{q\nu} - \Omega) \rangle_{kn} \rangle_{k'n'}. \quad (60)$$

Here g_S is the number of electron states per spin and atom at the Fermi level, M is the ion mass, $\omega_{q\nu}$ is the frequency of phonons with momentum $\mathbf{q} = \mathbf{k} - \mathbf{k}'$ and branch index ν , and the spin-flip matrix element is

$$g_{kn\uparrow, k'n'\downarrow}^\nu \equiv |\mathbf{u}_{q\nu} \cdot (\Psi_{kn\uparrow}, \nabla V \Psi_{k'n'\downarrow})|^2, \quad (61)$$

where $\mathbf{u}_{q\nu}$ is the polarization of the phonon with momentum \mathbf{q} and in branch ν . The brackets $\langle \dots \rangle_{kn}$ in Eq. (60) denote Fermi-surface averaging. The Bloch wave functions for this calculation are chosen to satisfy $(\Psi_{kn\uparrow}, \hat{\sigma}_z \Psi_{kn\uparrow}) = -(\Psi_{kn\downarrow}, \hat{\sigma}_z \Psi_{kn\downarrow})$. The periodic lattice-ion interaction V contains both scalar and spin-orbit parts: $V = V_{sc} + V_{so}$.

There are two important relations giving an order-of-magnitude estimate of τ_s , as well as its temperature dependence: the Elliott and the Yafet relations. The Elliott relation relates τ_s to the shift Δg of the electronic g factor from the free-electron value of $g_0 = 2.0023$ (Elliott, 1954):

$$1/\tau_s \approx (\Delta g)^2 / \tau_p, \quad (62)$$

where τ_p is the momentum relaxation time. This relation follows from the fact that for a momentum scattering interaction V_i the spin-flip scattering probability in the Born approximation is proportional to $|(\Psi_{kn\uparrow}, V_i \Psi_{k'n'\downarrow})|^2 \approx |b|^2 \times |(\Psi_{kn\uparrow}, V_i \Psi_{k'n'\downarrow})|^2$. Realizing that the spin-conserving scattering probability gives the momentum relaxation rate, after a Fermi-surface averaging we get the estimate

$$1/\tau_s \approx \langle b^2 \rangle / \tau_p. \quad (63)$$

On the other hand, Δg is determined by the expectation value of \hat{l}_z , the z-component of the orbital momentum in a Bloch state. Without the spin-orbit interaction this expectation value is zero. Considering the spin-orbit interaction to be a small parameter, we find by perturbation theory $\Delta g \approx |b|$, which combines with Eq. (63) to give the Elliott relation Eq. (62). An empirical test of the Elliott relation for simple metals when spin relaxation is due to thermal phonons (Beuneu and Monod, 1978) gives the revised estimate

$$1/\tau_s \approx 10 \times (\Delta g)^2 / \tau_p. \quad (64)$$

The Elliott relation is only a very rough estimate of τ_s . The experimentally relevant ratio τ_p/τ_s depends on the scattering mechanism. The ratio is different for scattering off impurities, boundaries, or phonons, although one would expect it to be within an order of magnitude. For example, scattering by heavy impurities induces an additional spin relaxation channel where spin flip is due to the spin-orbit interaction induced by the impurities. Equation (63) then does not hold. Scattering by phonons is too complex to be simply equated with the ratio τ_p/τ_s for impurity or boundary scattering. However, the ratio is comparable for scattering by light impurities and by boundaries. The ratio τ_p/τ_s for impurity and phonon scattering in Al and Cu is compared by Jedema *et al.* (2003).

The Yafet relation is only qualitative, connecting the temperature dependence of $1/T_1$ with that of the resistivity ρ :

$$1/T_1(T) \sim \langle b^2 \rangle \rho(T), \quad (65)$$

as follows directly from Eq. (63) after realizing that $1/\tau_p(T) \sim \rho(T)$. By careful symmetry considerations Yafet (1963) proved that $1/T_1 \sim T^5$ at low temperatures, similarly to the Bloch-Grüneisen law for resistivity, justifying Eq. (65) over a large temperature range. Yafet's T^5 law stems from the nontrivial fact that for spin-flip electron-phonon scattering $g_{kn\uparrow, k'n'\downarrow}^\nu \sim (\mathbf{k} - \mathbf{k}')^4$ as $\mathbf{k} \rightarrow \mathbf{k}'$ (Yafet, 1963), while only a quadratic dependence holds for the spin-conserving scattering. This corresponds to the long-wavelength behavior $\alpha_S^2 F(\Omega) \sim \Omega^4$ of the spin-flip Eliashberg function. The Yafet relation was tested experimentally by Monod and Beuneu (1979). This work led to a deeper understanding of spin relaxation processes in polyvalent metals (Fabian and Das Sarma, 1998; Silsbee and Beuneu, 1983).

Realistic calculations of the EY τ_s in semiconductors can be performed analytically using approximations of the band and phonon structures, as most important states are usually around high symmetry points. Here we give a formula for the spin relaxation of conduction electrons with energy $E_{\mathbf{k}}$ in the frequently studied case of III-V semiconductors (Chazalviel, 1975; Pikus and Titkov, 1984):

$$\frac{1}{\tau_s(E_{\mathbf{k}})} = A \left(\frac{\Delta_{so}}{E_g + \Delta_{so}} \right)^2 \left(\frac{E_{\mathbf{k}}}{E_g} \right)^2 \frac{1}{\tau_p(E_{\mathbf{k}})}, \quad (66)$$

where $\tau_p(E_{\mathbf{k}})$ is the momentum scattering time at energy $E_{\mathbf{k}}$, E_g is the energy gap, and Δ_{so} is the spin-orbit splitting of the valence band (see Fig. 6). The Numerical factor A , which is of order 1, depends on the dominant scattering mechanism (charge or neutral impurity, phonon, electron-hole). Analytic formulas for the EY mechanism due to electron-electron scattering are given by Boguslawski (1980).

Equation (66) shows that the EY mechanism is important for small-gap semiconductors with large spin-orbit

splitting (the prototypical example is InSb). For degenerate electron densities the spin relaxation time is given by Eq. (66) with $E_{\mathbf{k}} = E_F$, while for nondegenerate densities the thermal averaging leads to a substitution of thermal energy $k_B T$ for $E_{\mathbf{k}}$ and thermal-averaged momentum relaxation time for τ_p . To estimate τ_s from Eq. (66) one needs to know τ_p . It often suffices to know the doping or temperature dependence of τ_p to decide on the relevance of the EY mechanism (Pikus and Titkov, 1984).

The temperature dependence of $1/\tau_s$ for metals and degenerate semiconductors follows the dependence of $1/\tau_p$. In metals this means a constant at low T and a linear increase at large T . For nondegenerate semiconductors $1/\tau_s(T) \sim T^2/\tau_p(T)$. In the important case of scattering by charged impurities ($\tau_p \sim T^{3/2}$) $1/\tau_s \sim T^{1/2}$. The magnetic field dependence of the EY spin relaxation has not been systematically investigated. At low temperatures, where cyclotron effects become important, one needs to average over cyclotron trajectories on the Fermi surface need to obtain $1/\tau_s$. We expect that such averaging leads, in general, to an increase in $1/T_1$, especially in systems where spin hot spots are important (see Sec. III.C).

2. D'yakonov-Perel' mechanism

An efficient mechanism of spin relaxation due to spin-orbit coupling in systems lacking inversion symmetry was found by D'yakonov and Perel' (1971e). Without inversion symmetry the momentum states of the spin up and spin down electrons are not degenerate: $E_{\mathbf{k}\uparrow} \neq E_{\mathbf{k}\downarrow}$. Kramer's theorem still dictates that $E_{\mathbf{k}\uparrow} = E_{-\mathbf{k}\downarrow}$. Most prominent examples of materials without inversion symmetry come from groups III-V (such as GaAs) and II-VI (ZnSe) semiconductors, where inversion symmetry is broken by the presence of two distinct atoms in the Bravais lattice. Elemental semiconductors like Si possess inversion symmetry in the bulk, so the DP mechanism does not apply to them. In heterostructures the symmetry is broken by the presence of asymmetric confining potentials.

Spin splittings induced by inversion asymmetry can be described by introducing an intrinsic \mathbf{k} -dependent magnetic field $\mathbf{B}_i(\mathbf{k})$ around which electron spins precess with Larmor frequency $\Omega(\mathbf{k}) = (e/m)\mathbf{B}_i(\mathbf{k})$. The intrinsic field derives from the spin-orbit coupling in the band-structure. The corresponding Hamiltonian term describing the precession of electrons in the conduction band is

$$H(\mathbf{k}) = \frac{1}{2}\hbar\hat{\sigma} \cdot \Omega(\mathbf{k}), \quad (67)$$

where $\hat{\sigma}$ are the Pauli matrices. Momentum-dependent spin precession described by H , together with momentum scattering characterized by momentum relaxation

time τ_p ,⁷⁴ leads to spin dephasing. While the microscopic expression for $\Omega(\mathbf{k})$ needs to be obtained from the band structure, treating the effects of inversion asymmetry by introducing intrinsic precession helps to give a qualitative understanding of spin dephasing. It is important to note, however, that the analogy with real Larmor precession is not complete. An applied magnetic field induces a macroscopic spin polarization and magnetization, while H of Eq. (67) produces an equal number of spin up and spin down states.

Two limiting cases can be considered: (i) $\tau_p\Omega_{av} \gtrsim 1$ and (ii) $\tau_p\Omega_{av} \lesssim 1$, where Ω_{av} is an average magnitude of the intrinsic Larmor frequency $\Omega(\mathbf{k})$ over the actual momentum distribution. Case (i) corresponds to the situation in which individual electron spins precess a full cycle before being scattered to another momentum state. The total spin in this regime initially dephases reversibly due to the anisotropy in $\Omega(\mathbf{k})$. The spin dephasing rate,⁷⁵ which depends on the distribution of values of $\Omega(\mathbf{k})$, is in general proportional to the width $\Delta\Omega$ of the distribution: $1/\tau_s \approx \Delta\Omega$. The spin is irreversibly lost after time τ_p , when randomizing scattering takes place.

Case (ii) is what is usually meant by the D'yakonov-Perel' mechanism. This regime can be viewed from the point of view of individual electrons as a spin precession about fluctuating magnetic fields, whose magnitude and direction change randomly with the average time step of τ_p . The electron spin rotates about the intrinsic field at an angle $\delta\phi = \Omega_{av}\tau_p$, before experiencing another field and starting to rotate with a different speed and in a different direction. As a result, the spin phase follows a random walk: after time t , which amounts to t/τ_p steps of the random walk, the phase progresses by $\phi(t) \approx \delta\phi\sqrt{t/\tau_p}$. Defining τ_s as the time at which $\phi(t) = 1$, the usual motional narrowing result is obtained: $1/\tau_s = \Omega_{av}^2\tau_p$ (see footnote 70). The faster the momentum relaxation, the slower the spin dephasing. The difference between cases (i) and (ii) is that in case (ii) the electron spins form an ensemble that directly samples the distribution of $\Omega(\mathbf{k})$, while in case (ii) it is the distribution of the *sums* of the intrinsic Larmor frequencies (the total phase of a spin after many steps consists of a sum of randomly selected frequencies multiplied by τ_p), which, according to the central limit theorem, has a significantly reduced variance. Both limits (i) and (ii) and the transition between them have been experimentally demonstrated in n-GaAs/AlGaAs quantum wells by observing temporal spin oscillations over a large range of temperatures (and thus τ_p) (Brand *et al.*, 2002).

A more rigorous expression for τ_s in regime (ii) has been obtained by solving the kinetic rate equation for

⁷⁴ In the qualitative reasonings below we use τ_p instead of the effective correlation time $\tilde{\tau}$ for Ω during momentum scattering; $\tilde{\tau}$ is defined later, in Eq. (69).

⁷⁵ The reversible decay need not be exponential.

a spin-dependent density matrix (D'yakonov and Perel', 1971d,e). If the band structure is isotropic and scattering is both elastic and isotropic, evolution of the z-component of spin \mathbf{s} is (Pikus and Titkov, 1984)

$$\dot{s}_z = -\tilde{\tau}_l \left[s_z \overline{(\Omega^2 - \Omega_z^2)} - s_x \overline{\Omega_x \Omega_z} - s_y \overline{\Omega_y \Omega_z} \right], \quad (68)$$

where the bar denotes averaging over directions of \mathbf{k} . Analogous expressions for \dot{s}_x and \dot{s}_y can be written by index permutation. The effective momentum scattering time is introduced as

$$1/\tilde{\tau}_l = \int_{-1}^1 W(\theta) [1 - P_l(\cos \theta)] d\cos \theta, \quad (69)$$

where $W(\theta)$ is the rate of momentum scattering through angle θ at energy $E_{\mathbf{k}}$, and P_l is the Legendre polynomial whose order l is the power of \mathbf{k} in $\Omega(\mathbf{k})$. [It is assumed that $\Omega(\mathbf{k}) \sim k^l$ in Eq. (68).] In two dimensions $P_l(\cos \theta)$ is replaced by $\cos(l\theta)$ in Eq. (69) for the l th polar harmonic of $\Omega(\mathbf{k})$ (Pikus and Pikus, 1995). Since it is useful to express the results in terms of the known momentum relaxation times⁷⁶ $\tau_p = \tilde{\tau}_1$, one defines⁷⁷ $\gamma_l = \tau_p/\tilde{\tau}_l$ to measure the effectiveness of momentum scattering in randomizing Larmor frequencies; $\tilde{\tau}_l$ accounts for the relative angle between Ω before and after scattering. Generally $\gamma_l > 1$ for $l > 1$, that is, momentum scattering is more effective in randomizing spins than in randomizing momentum.

Comparing with the Bloch-Torrey equations (52)–(54), for $\mathbf{B} = \mathbf{0}$ and no spin diffusion, we see that spin decay is described by the tensor $1/\tau_{s,ij}$ (here i and j are the Cartesian coordinates) whose diagonal $1/\tau_{s,ii}$ and off-diagonal $1/\tau_{s,ij}$, for $i \neq j$, terms are

$$1/\tau_{s,ii} = \gamma_l^{-1} \tau_p (\overline{\Omega^2} - \overline{\Omega_i^2}), \quad 1/\tau_{s,ij} = -\gamma_l^{-1} \tau_p \overline{\Omega_i \Omega_j}. \quad (70)$$

In general, spin dephasing depends on the spin direction and on the dephasing rates of the perpendicular spin components. Equations (70) are valid for small magnetic fields, satisfying $\Omega_0 \tau_p \ll 1$, where Ω_0 is the Larmor frequency of the external field.

The most important difference between the EY and the DP mechanism is their opposite dependence on τ_p . While increased scattering intensity makes the EY mechanism more effective, it decreases the effectiveness of the the DP processes. In a sense the two mechanisms are similar to collision broadening and motional narrowing in NMR (Slichter, 1989). Indeed, in the EY process the precession frequency is conserved between collisions and the loss of

phase occurs only in the short time *during* collision. The more collisions there are, the greater is the loss of phase memory, in analogy with collision broadening of spectral lines. On the other hand, in DP spin dephasing, spin phases are randomized *between* collisions, since electrons precess with different frequencies depending on their momenta. Spin-independent collisions with impurities or phonons do not lead to phase randomization during the collision itself, but help to establish the random-walk-like evolution of the phase, leading to motional narrowing. While these two mechanisms coexist in systems lacking inversion symmetry, their relative strength depends on many factors. Perhaps the most robust trend is that the DP mechanism becomes more important with increasing band gap and increasing temperature.

In the rest of the section we apply Eq. (70) to the study of spin dephasing in bulk and two-dimensional III-V semiconductor heterostructures.

a. Bulk III-V semiconductors. In bulk III-V semiconductors the intrinsic Larmor frequency vector of Eq. (67) due to the lack of inversion symmetry is (D'yakonov and Perel', 1971d)

$$\Omega(\mathbf{k}) = \alpha \hbar^2 (2m_c^3 E_g)^{-1/2} \kappa, \quad (71)$$

where

$$\kappa = [k_x(k_y^2 - k_z^2), k_y(k_z^2 - k_x^2), k_z(k_x^2 - k_y^2)]. \quad (72)$$

Here k_i are the lattice wave-vector components along the crystal principal axes. The material-specific parameters are the band gap E_g and the conduction electron mass m_c ; α is a dimensionless parameter specifying the strength of the spin-orbit interaction. The spin splitting described by Eq. (71) is proportional to the cube of the lattice momentum, as was first found by Dresselhaus (1955). For GaAs $\alpha \approx 0.07$ (Marushchak *et al.*, 1984). Spin splitting of conduction and heavy and light holes in GaAs quantum wells, due to bulk inversion asymmetry was calculated by Rashba and Sherman (1988).

Substituting Eq. (71) for Ω in Eq. (70), and using $\overline{\kappa_i \kappa_j} = (4/105)k^6 \delta_{ij}$, we obtain the expected result that the off-diagonal elements of $1/\tau_{s,ij}$ vanish for cubic systems and the diagonal elements are all equal to (Pikus and Titkov, 1984)

$$1/\tau_s(E_{\mathbf{k}}) = \frac{32}{105} \gamma_3^{-1} \tau_p(E_{\mathbf{k}}) \alpha^2 \frac{E_{\mathbf{k}}^3}{\hbar^2 E_g}. \quad (73)$$

The above expression describes DP spin dephasing of degenerate ($E_{\mathbf{k}} = E_F$) or hot⁷⁸ electrons in bulk III-V semiconductors. For impurity scattering $\gamma_3 \approx 6$, for

⁷⁶ In fact, normal (not umklapp) electron-electron collisions should also be included in the effective spin randomization time $\tilde{\tau}$, though they do not contribute to the momentum relaxation time which appears in the measured mobility (Glazov and Ivchenko, 2002, 2003).

⁷⁷ Pikus and Titkov (1984) initially define γ_l as here, but later evaluate it, inconsistently, as the inverse $\gamma_l \rightarrow \gamma_l^{-1}$.

⁷⁸ This is strictly true only if the spin relaxation of the hot electrons is faster than energy relaxation by optical phonon emission, which is rarely the case. One has to consider either the spin re-

acoustic phonons $\gamma_3 \approx 1$, while for optical polar phonons $\gamma_3 \approx 41/6$. The temperature dependence of $1/\tau_s$ comes from the temperature dependence of τ_p . For the important case of charged impurity scattering ($\tau_p \sim T^{3/2}$), $1/\tau_s \sim T^{3/2}$.

Compared to the EY expression, Eq. (66), the DP spin dephasing increases much faster with increasing electron energy and is expected to be dominant at large donor doping levels and at high temperatures. The EY mechanism can be dominant in small-band-gap and large spin-orbit-splitting materials. The two mechanisms can also be easily distinguished by their opposite dependence on momentum relaxation. Contrary to the EY mechanism, greater impurity density will decrease the importance of the DP processes. The most frequently used ways to distinguish between various methods of spin relaxation are comparing the electron density (through the variation of the Fermi energy) and the temperature dependences of $1/\tau_s$ with the theoretical estimates. Since the prefactors may vary with different scattering mechanisms, it is best to deduce $\tau_p(E_{\mathbf{k}})$ and $\tau_p(T)$ from mobility measurements and use Eqs. (66) and (73) or the equations given below for $1/\tau_s(T)$.

Another interesting distinction between the two mechanisms is revealed by the dependence of their spin diffusion length $L_s = \sqrt{D\tau_s}$ on momentum scattering. Since $D \sim \tau_p$, for EY $L_s \sim \tau_p$, while for DP L_s does not depend on the momentum scattering time and for a degenerate electron system should be a constant independent of T , of the order of v_F/Ω_{av} . We do not know of an experimental verification of this distinction.

If the electrons obey nondegenerate statistics, which is the usual case of p-doped materials, thermal averaging over the Boltzmann distribution gives (Pikus and Titkov, 1984).

$$1/\tau_s = Q\tau_m\alpha^2 \frac{(k_B T)^3}{\hbar^2 E_g}, \quad (74)$$

where $\tau_m = \langle \tau_p(E_{\mathbf{k}})E_{\mathbf{k}} \rangle / \langle E_{\mathbf{k}} \rangle$. The coefficient Q , which is of order 1, is

$$Q = \frac{16}{35} \gamma_3^{-1} \left(\nu + \frac{7}{2} \right) \left(\nu + \frac{5}{2} \right), \quad (75)$$

where the power law $\tau_p \sim E_{\mathbf{k}}^\nu$ is assumed for momentum relaxation time. For scattering by ionized impurities $Q \approx 1.5$, while scattering by polar optical or piezoelectric phonons gives $Q \approx 0.8$, and scattering by acoustic phonons (deformation potential) $Q \approx 2.7$ (Pikus and

Titkov, 1984). The temperature behavior of DP spin dephasing in nondegenerate samples is $1/\tau_s \sim T^3\tau_m(T)$. For scattering by charged impurities $1/\tau_s \sim T^{9/2}$.

Application of longitudinal (to the initial spin direction) magnetic field suppresses the DP mechanism (Pikus and Titkov, 1984) for two reasons: (i) The B -field suppresses precession along the transverse intrinsic fluctuating fields when $\Omega_L\tau_p > 1$, where Ω_L is the Larmor precession due to B . (ii) $\Omega_{\mathbf{k}}$ is orbitally averaged, which has a similar effect to averaging by random scattering, when $\Omega_c\tau_p > 1$, where Ω_c is the cyclotron frequency. Since for conduction electrons $m_c \ll m_e$, it follows that $\Omega_c \gg \Omega_L$, the orbital motion induced by B is the cause for suppression of spin relaxation in semiconductors.

b. Two-dimensional III-V semiconductor systems. In two-dimensional III-V semiconductor systems (quantum wells and heterostructures) there are two distinct Hamiltonian terms that contribute to DP spin dephasing: the bulk inversion asymmetry term H_{BIA} and the structure inversion asymmetry term, H_{SIA} , which appears only in asymmetric systems. Both H_{BIA} and H_{SIA} lead to spin splitting of the conduction band linear in \mathbf{k} . The two terms, however, predict a different dependence of τ_s on the quantum-well orientation relative to the principal axes. Figure 15 shows the vector field patterns of the intrinsic magnetic fields for both bulk and spin inversion asymmetry.

The bulk inversion asymmetry term comes from the bulk Dresselhaus spin splitting, Eq. (71). Treating wave vectors \mathbf{k} in Eq. (71) as operators $\hat{\mathbf{k}} = -i\nabla$, and evaluating $\boldsymbol{\Omega}$ as the expectation value in the confined states, leads to momentum quantization along the confinement unit vector \mathbf{n} of the quantum well (QW). In the following, \mathbf{k} denotes the wave vector for a Bloch state propagating in the plane, and $k_n^2 \equiv \langle (\hat{\mathbf{k}} \cdot \mathbf{n})^2 \rangle$ denotes the expectation value of the square of the component of the wave-number operator normal to the plane in the lowest-subband state. For a rectangular QW of width a , $k_n^2 = (\pi/a)^2$. For a triangular well of confining potential eEz , $k_n^2 \approx 0.7794(2m_e E/\hbar^2)^{2/3}$ [see, for example, de Sousa and Das Sarma (2003b)]. Quantum averaging of κ can be done using the formula

$$\langle \hat{k}_i \hat{k}_j \hat{k}_l \rangle = k_n^2 (k_i n_j n_l + k_j n_l n_i + k_l n_i n_j) + k_i k_j k_l. \quad (76)$$

This readily gives (D'yakonov and Kachorovskii, 1986)

$$\kappa_x = k_n^2 [2n_x (n_y k_y - n_z k_z) + k_x (n_y^2 - n_z^2)], \quad (77)$$

and similarly for κ_y and κ_z by index permutation. Terms cubic in k were omitted from the above equation, assuming that for narrow QW's $k^2 \ll k_n^2$. The explicit knowledge of κ is useful in qualitative analysis of spin dephasing for particular orientations of QW's.

The spin dephasing tensor $1/\tau_{s,ij}$, defined in Eq. (70),

laxation at different energy levels during the cascade process of optical phonon emission or, if the optical phonon emission is particularly fast, spin relaxation only during the final stages of energy relaxation by acoustic phonon emission (see Pikus and Titkov (1984)).

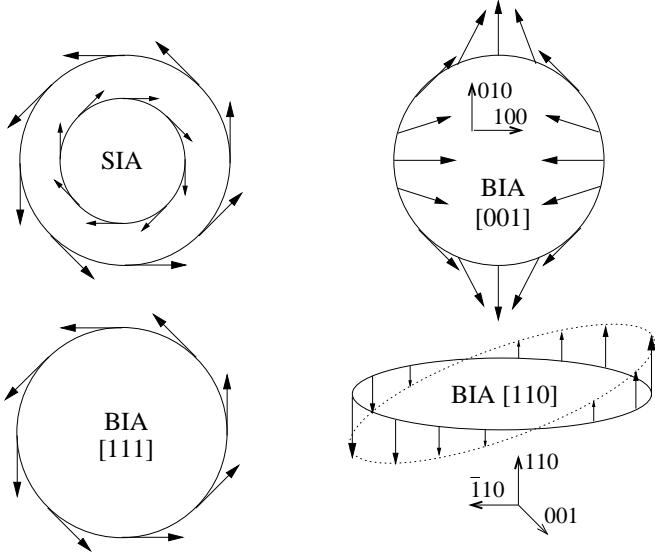


FIG. 15 Vector fields $\Omega(\mathbf{k}) \sim \kappa(\mathbf{k})$ on the Fermi surface (here a circle) for the structure inversion asymmetry (SIA) and bulk inversion asymmetry (BIA). Since $\Omega(\mathbf{k})$ is also the spin quantization axis, the vector pattern is also the pattern of the spin on the Fermi surface. As the opposite spins have different energies, the Fermi circle becomes two concentric circles with opposite spins. This is shown here only for the SIA case, but the analogy extends to all examples. The field for BIA [110] lies perpendicular to the plane, with the magnitude varying along the Fermi surface. All other cases have constant fields lying in the plane.

is readily evaluated using Eqs. (71) and (77)⁷⁹

$$1/\tau_{s,ij} = (\delta_{ij} \text{Tr}\hat{\nu} - \nu_{ij}) / \tau_s^0(E_{\mathbf{k}}), \quad (78)$$

where

$$\frac{1}{\tau_s^0(E_{\mathbf{k}})} = \frac{\alpha^2 \hbar^2 (k_n^2)^2}{2m_c^2 E_g} E_{\mathbf{k}} \tau_p(E_{\mathbf{k}}). \quad (79)$$

The tensor $\hat{\nu}$ depends on the orientation of \mathbf{n} with respect to the principal crystal axes (D'yakonov and Kachorovskii, 1986)⁸⁰

$$\nu_{xx} = 4n_x^2(n_y^2 + n_z^2) - (n_y^2 - n_z^2)^2(9n_x^2 - 1), \quad (80)$$

$$\nu_{xy} = n_x n_y [9(n_x^2 - n_z^2)(n_y^2 - n_z^2) - 2(1 - n_z^2)], \quad (81)$$

and analogously for other components.

We follow D'yakonov and Kachorovskii (1986) in discussing the three important cases of [001], [111], and [110] quantum wells. For [001],

$$\Omega(\mathbf{k}) \sim \kappa = k_n^2(-k_x, k_y, 0). \quad (82)$$

While the magnitude of $\Omega(\mathbf{k})$ is constant over the Fermi surface, the directions follow a “breathing” pattern as shown in Fig. 15. The spin relaxation times follow from Eq. (79): $1/\tau_{s,xx} = 1/\tau_{s,yy} = 1/2\tau_{zz} = 1/\tau_s^0$. Defining $1/\tau_{s,\parallel}$ and $1/\tau_{s,\perp}$ as spin dephasing rates of spins parallel and perpendicular to the plane, one obtains

$$1/\tau_{s,\parallel} = 1/2\tau_{s,\perp} = 1/\tau_s^0. \quad (83)$$

As expected for the case of the in-plane field, the lifetime of a spin parallel to the plane is twice that of a spin perpendicular to the plane.

For [111] QW's,

$$\Omega(\mathbf{k}) \sim \kappa = 2/\sqrt{3}k_n^2(\mathbf{k} \times \mathbf{n}). \quad (84)$$

The intrinsic magnetic field lies in the plane, having a constant magnitude (refer to Fig. 15). Spin relaxation rates are now $1/\tau_{s,ii} = 16/9\tau_s^0$ and $1/\tau_{s,i \neq j} = 4/9\tau_s^0$. By diagonalizing $1/\tau_{ij}$ we obtain

$$1/\tau_{s,\parallel} = 1/2\tau_{s,\perp} = 4/3\tau_s^0. \quad (85)$$

As for the [001] case, a perpendicular spin dephases twice as fast as a parallel one, since $\Omega(\mathbf{k})$ lies in the plane.

The most interesting case is the [110] orientation for which $1/\tau_{s,xx} = 1/\tau_{s,yy} = 1/2\tau_{s,zz} = -1/\tau_{s,xy} = 1/8\tau_s^0$. Other off-diagonal components vanish. Diagonalizing the tensor gives

$$1/\tau_{s,\parallel} = 1/4\tau_s^0, \quad 1/\tau_{s,\perp} = 0. \quad (86)$$

The perpendicular spin does not dephase. This is due to the fact that κ , unlike the previous cases, is always normal to the plane (see Fig. 15), and thus cannot affect the precession of the perpendicular spin. Indeed,

$$\Omega(\mathbf{k}) \sim \kappa = k_n^2(k_x/2)(-1, -1, 0), \quad (87)$$

where it is used that $\mathbf{k} \cdot \mathbf{n} = 0$. Spin dephasing in [110] QW's can still be due to the cubic terms in k left out of Eq. (77) or to other spin relaxation mechanisms. Note that the magnitude of $\Omega(\mathbf{k})$ changes along the Fermi surface. Electrons moving along [001] experience little spin dephasing.

The structure inversion asymmetry term arises from the Bychkov-Rashba spin splitting (Bychkov and Rashba, 1984a,b; Rashba, 1960) occurring in asymmetric QW's or in deformed bulk systems. The corresponding Hamiltonian is that of Eq. (67), with the precession vector

$$\Omega(\mathbf{k}) = 2\alpha_{BR}(\mathbf{k} \times \mathbf{n}). \quad (88)$$

Here α_{BR} is a parameter depending on spin-orbit coupling and the asymmetry of the confining electrostatic potentials arising from the growth process of the heterostructure. The splitting can also arise in nominally symmetric heterostructures with fluctuations in doping density (Sherman, 2003b). The Bychkov-Rashba field

⁷⁹ Averaging over the directions of \mathbf{k} in a plane perpendicular to \mathbf{n} can be performed by using $\overline{k_i k_j} = (k^2/2)(\delta_{ij} - n_i n_j)$.

⁸⁰ A trivial typo in ν_{xx} is corrected.

always lies in the plane, having a constant magnitude. As for the bulk inversion asymmetry case, the structure inversion asymmetry leads to a splitting of the Fermi surface, according to the direction of the spin pattern—parallel or antiparallel to $\Omega(\mathbf{k})$, as shown in Fig. 15. Perhaps the most appealing fact about structure inversion asymmetry is that α_{BR} can be tuned electrostatically, potentially providing an effective spin precession control without the need for magnetic fields (Levitov and Rashba, 2003; Rashba and Efros, 2003). This has led to one of the pioneering spintronic proposals by Datta and Das (1990) (see Sec. IV.E.1). Note that for the [111] orientation the bulk and structure inversion asymmetry terms have the same form.

Using the same procedure as for bulk inversion asymmetry, we describe the spin relaxation rate by Eq. (78) as

$$1/\tau_s^0 = 4\alpha_{BR}^2 \frac{m_c}{\hbar^2} E_k \tau_p \quad (89)$$

and

$$\nu_{ij} = 1 - n_i n_j. \quad (90)$$

Since the intrinsic precession vector $\sim \Omega(\mathbf{k})$ for the structure inversion asymmetry always lies in the plane, a perpendicular spin should dephase twice as fast as a spin in the plane. Indeed, by diagonalizing $1/\tau_{s,ij}$ one finds that

$$1/\tau_{s,\parallel} = 1/2\tau_{s,\perp} = 1/\tau_s^0 \quad (91)$$

holds for all QW orientations \mathbf{n} . This interesting fact qualitatively distinguishes structure from bulk inversion asymmetry and can be used in assessing the relative importance of the Dresselhaus and Rashba spin splittings in III-V heterostructure systems. If bulk and structure inversion asymmetry are of similar importance, the interference terms from the cross product $\bar{\Omega}_{BIA}\bar{\Omega}_{SIA}$ can lead to spin dephasing anisotropies within the plane, as was shown for [001] QW's in (Averkiev and Golub, 1999; Kainz *et al.*, 2003). This plane anisotropy can be easily seen by adding the corresponding vector fields in Fig. 15. Another interesting feature of bulk and structure inversion asymmetry fields is that injection of electrons along a quasi-one-dimensional channel can lead to large relaxation times for spins oriented along $\Omega(\mathbf{k})$, where \mathbf{k} is the wave vector for the states in the channel (Hammar and Johnson, 2002).

Model spin dephasing calculations based on structure inversion asymmetry were carried out by Pareek and Bruno (2002). Calculations of τ_s based on the DP mechanism, with structural asymmetry due to doping fluctuations in the heterostructure interface were performed by Sherman (2003b).

Research on spin inversion asymmetry is largely motivated by Datta-Das spin field-effect transistor proposal (see Secs. I.A and IV.E.1) in which α_{BR} is tailored by a gate. This tailoring, however, has been controversial and the microscopic origin of the Bychkov-Rashba Hamiltonian, and thus the interpretation of experimental results

on splitting in semiconductor heterostructures, has been debated. The Bychkov-Rashba Hamiltonian is often interpreted as arising from the electric field of the confining potential, assisted by external bias, which acts on a moving electron in a transverse direction. The relativistic transformation then gives rise to a magnetic field (spin-orbit coupling) acting on the electron spin. The parameter α_{BR} is then assumed to be directly proportional to the confining electrical field. This is in general wrong, since the average electric force acting on a confined particle of uniform effective mass is zero.

The asymmetry that gives rise to structure inversion asymmetry is the asymmetry in the band structure (including spin-orbit coupling) parameters of a heterostructure, such as the effective mass, or the asymmetry in the penetration of the electron wave function into the barriers (de Andrade e Silva *et al.*, 1997). The difficulty in understanding the influence of the external gates is caused by the lack of the understanding of the influence of the gate field on the asymmetry of the well. For a clear qualitative explanation of the involved physics see Pfeffer and Zawadzki (1999) and Zawadzki and Pfeffer (2001). Band-structure $k \cdot p$ calculations of α_{BR} for quantum wells in GaAs/AlGaAs heterostructures can be found in Pfeffer and Zawadzki (1995), Pfeffer (1997), Wissinger *et al.* (1998), and Kainz *et al.* (2003); a calculation for InGaAs/InP quantum wells is reported in Engels *et al.* (1997) and Schäpers *et al.* (1998), in InSb/InAlSb asymmetric quantum well it can be found in Pfeffer and Zawadzki (2003), and in p-InAs metal-oxide-semiconductor field-effect transistor channel in Lamari (2003). Adding to the controversy, Majewski and Vogl (2003) have recently calculated the structure inversion asymmetry by local density functional methods and concluded that the induced spin splitting arises from microscopic electric fields in asymmetric atomic arrangements at the interfaces, so that a large Bychkov-Rashba term can be present in otherwise symmetric quantum wells with no common atom.

Interpretation of experimental data on structure inversion asymmetry is difficult, especially at determining the zero magnetic field spin splitting (usually seen in Shubnikov-de Haas oscillations), which is masked by Zeeman splitting at finite fields. In addition, the splittings are small, typically less than 1 meV. The Bychkov-Rashba parameter was measured in GaSb/InAs/GaSb quantum wells ($\alpha_{BR} \approx 0.9 \times 10^{-11}$ eV m for 75 Å thick well) by Luo *et al.* (1990); in InAlAs/InGaAs/InAlAs quantum wells (20 nm), where also the gate voltage is obtained: α_{BR} ranged from 10^{-11} eV m at the depleting voltage of -1 V, to 5×10^{-12} eV m at +1.5 V. Weak antilocalization studies of InAlAs/InGaAs/InAlAs quantum wells have recently been used to study electron density dependence of α_{BR} by Koga *et al.* (2002a). Gate dependence of α_{BR} was also measured in modulation-doped InP/InGaAs/InP quantum wells (Engels *et al.*, 1997; Schäpers *et al.*, 1998). The observed values are several 10^{-12} eV m. On the other hand, there are ex-

perimental reports that either fail to observe the expected spin splitting due to Bychkov-Rashba field, or interpret the splitting differently [see, for example, Brosig *et al.* (1999) and Rowe *et al.* (2001)]. Furthermore, measurements of Heida *et al.* (1998) show a constant $\alpha_{BR} \approx 0.6 \times 10^{-11}$ eV m, independent of gate voltage, in asymmetric AlSb/InAs/AlSb quantum wells, demonstrating that control of α_{BR} may be difficult. In order to unify the different views on what exactly the Bychkov-Rashba spin splitting means and how the spin splitting can be tuned with gate voltage, more experimental efforts need to be devoted to this interesting topic.

3. Bir-Aronov-Pikus Mechanism

Spin relaxation of conduction electrons in p-doped semiconductors can also proceed through scattering, accompanied by spin exchange, by holes, as was first shown by Bir *et al.* (1975).

The exchange interaction between electrons and holes is governed by the Hamiltonian

$$H = AS \cdot \mathbf{J}\delta(\mathbf{r}), \quad (92)$$

where A is proportional to the exchange integral between the conduction and valence states, \mathbf{J} is the angular momentum operator for holes, \mathbf{S} is the electron spin operator, and \mathbf{r} is the relative position of electrons and holes.

The spin-flip scattering probability depends on the state of the holes (degenerate or nondegenerate, bound on acceptors or free, fast or slow). We present below the most frequently used formulas when assessing the relevance of the BAP mechanism. The formulas are valid for the usual cases of heavy holes $m_v \gg m_c$. For electron spin relaxation due to exchange with nondegenerate holes,

$$\frac{1}{\tau_s} = \frac{2}{\tau_0} N_a a_B^3 \frac{v_k}{v_B} \left[\frac{p}{N_a} |\psi(0)|^4 + \frac{5}{3} \frac{N_a - p}{N_a} \right], \quad (93)$$

where a_B is the exciton Bohr radius $a_B = \hbar^2 \epsilon / e^2 m_c$, p is the density of free holes, τ_0 is an exchange splitting parameter: $\hbar/\tau_0 = (3\pi/64)\Delta_{ex}/E_B$ (with E_B denoting the Bohr exciton energy, $E_B = \hbar^2/2m_c a_B^2$ and Δ_{ex} the exchange splitting of the excitonic ground state), and $v_B = \hbar/m_c a_B$; $|\psi(0)|^2$ is Sommerfeld's factor, which enhances the free hole contribution. For an unscreened Coulomb potential

$$|\psi(0)|^2 = \frac{2\pi}{\kappa} \left[1 - \exp \left(-\frac{2\pi}{\kappa} \right) \right]^{-1}, \quad (94)$$

where $\kappa = E_k/E_B$. For a completely screened potential $|\psi(0)|^2 = 1$.

If holes are degenerate and the electrons' velocity v_k is greater than the Fermi velocity of the holes', then

$$\frac{1}{\tau_s} = \frac{3}{\tau_0} p a_B^3 \frac{v_k}{v_B} \frac{k_B T}{E_{Fh}}, \quad (95)$$

where E_{Fh} is the hole Fermi energy. For degenerate holes $|\psi(0)|^2$ is of order 1. If electrons are thermalized, v_k needs to be replaced by the thermal velocity $v_e = (3k_B T/m_c)^{1/2}$.

The temperature dependence of τ_s is dominated by the temperature dependence of $|\psi(0)|^2$ as well as by p . The dependence on the acceptor density is essentially $1/\tau_s \sim N_a$ for nondegenerate/bound holes from Eq. (93) and $1/\tau_s \sim N_a^{1/3}$ for degenerate holes from Eq. (95). In between, $1/\tau_s$ is only weakly dependent on N_a . For GaAs $a_B \approx 114$ Å, $E_B \approx 4.9$ meV, $v_B \approx 1.7 \times 10^7$ cm·s⁻¹, $\tau_0 \approx 1 \times 10^{-8}$ s, and $\Delta_{ex} \approx 4.7 \times 10^{-5}$ eV (Aronov *et al.*, 1983).

The BAP mechanism coexists with the the EY and DP mechanisms in p-doped materials lacking inversion symmetry. The three mechanisms can be distinguished by their unique density and temperature dependences. A general trend is that the BAP dominates in heavily doped samples at small temperatures. At large temperatures even for large acceptor densities, the DP mechanism can become more important, due to its increased importance at large electron energies. Specific examples of the domain of importance for the three mechanisms are discussed in Sec. III.D.1. Model calculations of BAP processes for electrons in p-doped bulk and quantum wells were performed by Maialle (1996); Maialle and Degani (1997).

Another potentially relevant mechanism for spin relaxation of donor-bound electrons in p-doped semiconductors is the exchange interaction with holes bound to acceptors (D'yakonov and Perel', 1973b). The exchange interaction provides an effective magnetic field for electron spins to precess, leading to inhomogeneous dephasing. Both electron hopping and hole spin-flip motionally narrow the precession.

4. Hyperfine-interaction mechanism

The hyperfine interaction, which is the magnetic interaction between the magnetic moments of electrons and nuclei, provides an important mechanism (D'yakonov and Perel', 1973b) for ensemble spin dephasing and single-spin decoherence of localized electrons, such as those confined in quantum dots (QD) or bound on donors. The interaction is too weak to cause effective spin relaxation of free electrons in metals or in bulk semiconductors (Overhauser, 1953a), as it is strongly dynamically narrowed by the itinerant nature of electrons (see Sec. III.A.1). In addition to spin dephasing, the hyperfine interaction is relevant for spintronics as a means to couple, in a controlled way, electron and nuclear spins (D'yakonov and Perel', 1984a).

Localized electrons are typically spread over many lattice sites (10^4 – 10^6), experiencing the combined magnetic moments of many nuclei. In GaAs all the lattice nuclei carry the magnetic moment of 3/2 spin, while in Si the most abundant isotope, ²⁸Si, carries no spin and the

hyperfine interaction is due to ^{29}Si (natural abundance 4.67%) or the frequent donor ^{31}P , both of nuclear spin 1/2. As a result, an electron bound on a shallow donor in Si experiences only around 100 magnetic nuclei, and the effects of the hyperfine interaction are considerably smaller than in GaAs.

The effective Hamiltonian for the hyperfine interaction is the Fermi contact potential energy (Slrichter, 1989)

$$H = \frac{8\pi}{3} \frac{\mu_0}{4\pi} g_0 \mu_B \sum_i \hbar \gamma_{n,i} \mathbf{S} \cdot \mathbf{I}_i \delta(\mathbf{r} - \mathbf{R}_i), \quad (96)$$

where μ_0 is the vacuum permeability, $g_0 = 2.0023$ is the free-electron g factor, μ_B is the Bohr magneton, i is the label for nuclei at positions \mathbf{R}_i , \mathbf{S} , and \mathbf{I}_i are, respectively, electron and nuclear spin operators expressed in the units of \hbar , and $\gamma_{n,i}$ is the nuclear gyromagnetic ratio. We stress that it is the electron g factor g_0 and not the effective g that appears in the hyperfine interaction, Eq. (96), as shown by Yafet (1961) [see also Paget *et al.* (1977)]. It follows that the spin of an electron in an orbital state $\psi(\mathbf{r})$ experiences magnetic field

$$\mathbf{B}_n = \frac{2\mu_0}{3} \frac{g_0}{g} \sum_i \hbar \gamma_{n,i} \mathbf{I}_i |\psi(\mathbf{R}_i)|^2, \quad (97)$$

where g is the effective g factor of the electron. The electron Zeeman splitting due to the average B_n corresponds to a field of ~ 1 T or thermal energy of 1 K, for a complete nuclear polarization (Paget *et al.*, 1977).

There are three important regimes in which the hyperfine interaction leads to spin dephasing of localized electrons:

(i) In the limit of small orbital and spin correlation between separated electron states and nuclear spin states, spatial variations in \mathbf{B}_n lead to inhomogeneous dephasing of the spin ensemble, with the rate proportional to the r.m.s. of B_n , given by the corresponding thermal or nonequilibrium distribution of the nuclear spins. Such inhomogeneous dephasing is seen by electron spin resonance (ESR) experiments on donor states both in Si (Feher and Gere, 1959) and in GaAs (Seck *et al.*, 1997). This effect can be removed by spin echo experiments (in Si donor states performed, for example, by Gordon and Browers (1958)). The spread in the Larmor precession period due to the variance in B_n in GaAs is estimated to be around 1 ns (Dzhioev *et al.*, 2002b; Merkulov *et al.*, 2002)).

(ii) Temporal fluctuations in B_n , which can occur due to nuclear dipole-dipole interactions, lead to irreversible spin dephasing and decoherence of electron spins. Such processes are sometimes referred to as spectral diffusion, since the electron Zeeman levels split by B_n undergo random shifts (de Sousa and Das Sarma, 2003c). The typical time scale for the fluctuations in GaAs is given by the nuclear Larmor precession period in the field of neighboring nuclei and is of order 100 μs (Merkulov *et al.*, 2002). Nuclear moments also precess (and orient) in the magnetic

fields of polarized electrons, an effect important in optical orientation (Meier and Zakharchenya (Eds.), 1984), where the feedback from this precession can be directly observed through the modulated precession of electron spins. The time scale for the Larmor precession of nuclear spins in hyperfine fields is 1 μs in GaAs (Merkulov *et al.*, 2002), so this effect does not lead to motional narrowing of B_n ; electron spins precess many times before the nuclear spin flips.

(iii) In the presence of strong orbital correlations (electron hopping or recombination with acceptor hole states) or spin (direct exchange interaction) between neighboring electron states, spin precession due to B_n is motionally narrowed. While the direct spin exchange interaction does not cause ensemble spin relaxation (the total spin is preserved in spin flip-flops), it leads to individual spin decoherence, which can be much faster than what is inferred from T_2 . This effect is much more pronounced in GaAs than in Si, since the donor states spread to greater distances, and thus even in the low-doping limits ($\approx 10^{14} \text{ cm}^{-3}$ donors) the exchange interaction can be rather large, masking the effects of temporal fluctuations of B_n (see Sec. III.D.3). Many useful parameters for evaluating effective magnetic fields and precession frequencies due to the HFI mechanism in GaAs are given by Paget *et al.* (1977).

Ensemble spin dephasing due to the HFI mechanism in an external magnetic field has been studied by D'yakonov and Perel' (1973b), who found suppression of $1/\tau_s$ if the external field is greater than B_n for regime (i), or a smaller Larmor precession period than the correlation time for random changes in B_n , in regime (iii). due to the external field.

Calculations of τ_s using the HFI mechanism were performed for shallow donor states in Si at low temperatures and magnetic fields (Saikin *et al.*, 2002), for electron spins in QD's (Merkulov *et al.*, 2002; Semenov and Kim, 2003), and even for the case of conduction electrons in semiconductors (revisited by Pershin and Privman (2003c)). Spin relaxation processes due to phonon-assisted HFI were investigated in GaAs QD's, but were found to be ineffective (Erlingsson *et al.*, 2001). Unfortunately, there are still too few experimental data to make conclusions about the merits of specific models of the HFI mechanism.

Spin decoherence times for single-electron spins were recently computed for case (ii) by Khaetskii *et al.* (2002, 2003), who studied spin coherence time τ_{sc} of a single electron spin in the regime in which the electron Larmor period due to B_n is much shorter than the correlation time of the nuclear magnetic field fluctuations. Realistic estimates of HFI spin dephasing in GaAs QD's were given by de Sousa and Das Sarma (2003a,c), who offer reasons why mechanism (ii) should dominate spin decoherence in GaAs QD's of radius smaller than 100 nm. For instance, in a 50 nm wide QD the estimated τ_{sc} is $\approx 50 \mu\text{s}$, large enough for quantum computing applications (see Sec. IV.F.). This claim is supported by the recent mea-

surement of the spin dephasing time of about 60 ms of an isolated spin in a phosphorus donor in isotopically pure ^{28}Si , by spin echo measurements (Tyrlyshkin *et al.*, 2003).

C. Spin relaxation in metals

The spin relaxation time of conduction electrons in metals has been measured by both CESR and spin injection techniques. Typical values of τ_s were found to be 0.1 to 1 ns, but the range of observed values is large, from pico- to microseconds. To our knowledge the longest τ_s reported for a metal—a microsecond—was found in high-purity sodium below 10 K (Kolbe, 1971).

The majority of simple metals are believed to follow the EY mechanism of spin relaxation, with the possible exception of Li (Damay and Sienko, 1976). This is supported by several facts:

(i) The EY processes give the right order of magnitude for τ_s (Elliott, 1954; Yafet, 1963), while other known possible spin relaxation mechanisms lead to much greater τ_s than what is observed (Overhauser, 1953a).

(ii) The temperature dependence of τ_s is consistent with the EY mechanism: $1/\tau_s$ is constant at low temperatures, indicating impurity spin-flip scattering, while at high temperatures $1/\tau_s$ grows linearly with increasing T , consistent with phonon-induced spin relaxation.

(iii) The Elliott relationship, Eq. (62), has been tested for many important metals and found to be valid over many orders of magnitude of Δg (Beuneu and Monod, 1978) (this reference contains a useful collection of data for Δg). For the majority of metals tested (alkali and noble), a best fit gives the quantitative formula, Eq. (64) (Beuneu and Monod, 1978). (iv) The Yafet relation, Eq. (65), is satisfied for most metals with the known temperature dependence of τ_s (Fabian and Das Sarma, 1999c; Monod and Beuneu, 1979). The initially suggested deviation from the Yafet relation for several polyvalent metals (Al, Pd, Be, and Mg) was later resolved by spin-hot-spot theory (Fabian and Das Sarma, 1998, 1999c; Silsbee and Beuneu, 1983), to be described below. This work showed that the magnitudes of the spin-mixing probabilities b^2 , taken from atomic physics to test Eq. (65), should not be used in the solid-state environment. Various band-structure anomalies (spin hot spots), such as crossing of the Brillouin-zone boundaries, accidental degeneracy points, or symmetry points on the Fermi surface, can increase the atomic-physics-derived b^2 by several orders of magnitude, strongly enhancing spin relaxation in polyvalent metals as compared to simple estimates (Fabian and Das Sarma, 1998, 1999a).

(v) A realistic, first-principles calculation for Al (Fabian and Das Sarma, 1999b) (see Sec. III.C) using Eq. (59) shows excellent agreement with experiment.

CESR measurements of τ_s in various metals are nu-

merous.⁸¹

The spin injection technique (see Sec. II) was also used to measure τ_s for various metals, including Al (Jedema *et al.*, 2002a,b, 2003; Johnson and Silsbee, 1985), Au (Elezzabi *et al.*, 1996), Cu (Jedema *et al.*, 2001, 2003), and Nb (Johnson, 1994). In addition to CESR and spin injection, information about spin-orbit scattering times τ_{so} (see below) in various (but mostly noble) metals at low temperatures has been also obtained from weak localization magnetoresistance measurements on thin films (Bergmann, 1982) and tunneling spectroscopy of metallic nanoparticles (Petta and Ralph, 2001). Surface-scattering spin relaxation times in normal metals and superconductors are collected by Meservey and Tedrow (1978). Interesting results were obtained by injecting spin into superconductors. Using $\text{YBa}_2\text{Cu}_3\text{O}_{7-\delta}$, for example, data were interpreted (Fu *et al.*, 2002) to infer that the in-plane spin relaxation time is unusually long, about 100 μs at low temperatures to 1 μs close to the superconducting transition temperature. For quasiparticles moving along the c-axis, τ_s is more likely to be the usual spin-orbit-induced spin relaxation time, having the values of 10–100 ps. The microscopic origin of quasiparticle spin relaxation in cuprate superconductors is not yet known.

There is one more important time scale, the spin-orbit scattering time τ_{so} , that is often invoked in mesoscopic transport as a characteristic of spin relaxation processes. We discuss it briefly in connection to τ_s . The spin-orbit scattering time is the scattering time of Bloch electrons by the spin-orbit potential induced by impurities. The spin-orbit part of the Fourier transform of the impurity potential can be written as $ic(\mathbf{k} - \mathbf{k}')(\mathbf{k} \times \mathbf{k}') \cdot \sigma$, where $c(\mathbf{q})$ is proportional to the Fourier transform of the impurity potential. The spin-orbit scattering time then is (Werthamer, 1969)

$$1/\tau_{so} = \frac{2\pi}{\hbar} N_i \langle |c(\mathbf{k} - \mathbf{k}')|^2 |\mathbf{k} \times \mathbf{k}'| \rangle^2 \mathcal{N}(E_F), \quad (98)$$

where N_i is the impurity concentration, $\mathcal{N}(E_F)$ is the density of states per spin at the Fermi level, and the angle brackets denote Fermi-surface averaging. As a parameter τ_{so} also includes the spin-orbit coupling of the host lattice, in the sense of the EY mechanism. Note,

⁸¹ A list of selected metal includes Li (Damay and Sienko, 1976; Feher and Kip, 1955; Orchard-Webb *et al.*, 1970); Na (Feher and Kip, 1955; Kolbe, 1971; Vescil *et al.*, 1964), K (Walsh *et al.*, 1966a); Rb (Schultz and Shanabarger, 1966; Walsh *et al.*, 1966b); Cs (Schultz and Shanabarger, 1966; Walsh *et al.*, 1966b); Be (Cousins and Dupree, 1965; Orchard-Webb *et al.*, 1970), Mg (Bowring *et al.*, 1971); Cu (Lubzens and Schultz, 1976a; Monod and Schultz, 1982; Schultz and Latham, 1965); Au (Monod and Jánossy, 1977); Zn (Stesmans and Witters, 1981); Al (Lubzens and Schultz, 1976b); graphite (Matsubara *et al.*, 1991; Wagoner, 1960); Rb_3C_60 (Jánossy *et al.*, 1993); MgB_2 (Simon *et al.*, 2001). Various data on CESR τ_s are collected in (Beuneu and Monod, 1978; Monod and Beuneu, 1979).

however, that $\tau_s \neq \tau_{so}$, even at low temperatures where the impurity scattering dominates spin relaxation, since the spin-orbit scattering includes both spin-flip and spin-conserving processes, which, for isotropic scattering rates are in the ratio 2:1. In addition, the spin relaxation rate is twice the spin-flip scattering rate, since each spin flip equilibrates both spins equally. For isotropic systems $1/\tau_s \approx 4/(3\tau_{so})$. For a discussion of the effects of the DP processes on weak localization, see Knap *et al.* (1996).

We illustrate spin relaxation in metals on the case of Al, whose τ_s was measured by CESR and spin injection, and numerically calculated from first principles. The case is instructive since it illustrates both the general principles of the EY mechanism as well as the predicting power of, and the need for realistic band-structure calculations of τ_s .

Spin relaxation in Al was originally observed in CESR experiments, in which τ_s was measured at low temperatures, from 1 to 100 K (Lubzens and Schultz, 1976b). The spin relaxation rate $1/\tau_s$ was found to be independent of temperature below 10-20 K; at higher temperatures $1/\tau_s$ increases linearly with increasing T . The same behavior was later observed in the original spin injection experiment (Johnson and Silsbee, 1985, 1988d). Recently, τ_s was measured by spin injection at room temperature (Jedema *et al.*, 2002a,b, 2003). Unlike the CESR and the original spin injection experiments, which were performed on bulk samples, the room-temperature measurement used thin Al films, observing strong spin relaxation due to surface scattering.

Spin relaxation in Al, as well as in other polyvalent metals, at first appeared (Monod and Beuneu, 1979), in that a simple application of the Yafet relation, Eq. (65), yielded estimates for $1/\tau_s$ two orders of magnitude smaller than the observed data. Consider Na as a reference. The atomic $\lambda_{so}/\Delta E$ [cf. Eq. (58)] for Na and Al are within about 10% of each other (Beuneu and Monod, 1978; Monod and Beuneu, 1979), yet the corresponding τ_s for Al is about two orders of magnitude smaller than that for Na (Feher and Kip, 1955; Vescial *et al.*, 1964). This anomaly extends to the g factors as well. For Na, $\Delta g_{Na} \approx -8 \times 10^{-4}$ and for Al it is six times greater, $\Delta g_{Al} \approx -5 \times 10^{-3}$, while one would expect them to differ also by about 10%. Note, however, that the Elliott relation, Eq. (62), is unaffected by this discrepancy, as it predicts that $\tau_s(\text{Na})/\tau_s(\text{Al}) \approx 40$. It was later suggested (Silsbee and Beuneu, 1983) that this is due to accidental degeneracies in the two-band Fermi surface of Al.

A full theoretical description, supported by first-principles calculations, of spin relaxation in Al and other polyvalent metals led to the spin-hot-spots theory (Fabian and Das Sarma, 1998, 1999a,b,c). Spin hot spots are states on the Fermi surface that have anomalously large spin mixing probabilities $|b|^2 \approx (\lambda_{so}/\Delta E)^2$, arising from small energy gaps ΔE . Quite generally, such states occur near Brillouin-zone boundaries and accidental degeneracy points, or at high-symmetry points. The

condition for a spin hot spot is both a small band gap ΔE and nonvanishing λ_{so} .⁸²

If an electron hops in or out of a spin hot spot, the chance of spin flip dramatically increases. Although the total area of spin hot spots on the Fermi surface is small, their contribution to $1/\tau_s$ is dominant, due to the large value of their $|b|^2$ in the Fermi surface average $\langle |b|^2 \rangle$, as was shown by analytical arguments (Fabian and Das Sarma, 1998, 1999a). A realistic numerical calculation (Fabian and Das Sarma, 1999b) for Al, also showed that both the accidental degeneracies considered by (Silsbee and Beuneu, 1983) and states close to the Brillouin-zone boundaries dominate spin relaxation.

A realistic calculation of τ_s in Al, based on pseudopotentials and a realistic phonon description, has been performed by Fabian and Das Sarma (1999b) and compared to the experimental data available for $T < 100$ K (Johnson and Silsbee, 1985, 1988d). Figure 16 shows both the experiment and the theory. In the experimental data only the phonon contribution to $1/\tau_s$ is retained (Johnson and Silsbee, 1985); the constant background impurity scattering is removed. The figure shows a rapid decrease of τ_s with increasing T at low T , where the agreement between experiment and theory is very good. Above 200 K (the Debye temperature $T_D \approx 400$ K) the calculation predicts a linear dependence $\tau_s[\text{ns}] \approx 24 \times T^{-1}[\text{K}^{-1}]$. In the phonon-dominated linear regime the EY mechanism predicts that the ratio $a^{ph} = \tau_p/\tau_s$ does not depend on T (see Sec. III.B.1). The calculated value is $a_{th}^{ph} = 1.2 \times 10^{-4}$ (Fabian and Das Sarma, 1999b), showing that 10^4 phonon scatterings are needed to randomize electron spin.

An important step towards extending spin injection capabilities was undertaken recently by achieving spin injection into Cu and Al at room temperature (Jedema *et al.*, 2002a, 2001, 2002b, 2003); the measured data are unique in providing reliable values for spin diffusion lengths and spin relaxation times in these two important metals at room temperature. The measured values for Al are somewhat sensitive to the experimental procedure/data analysis: $\tau_s = 85$ ps (Jedema *et al.*, 2002b) and $\tau_s = 124$ ps (Jedema *et al.*, 2003), as compared to $\tau_s = 90$ ps predicted by the theory at $T = 293$ K. The room temperature experimental data are included in Fig. 16 for comparison. They nicely confirm the theoretical prediction. Less sensitive to data analysis is the ratio a^{ph} , for which the experiments give 1.1×10^{-4} (Jedema *et al.*, 2002b) and 1.3×10^{-4} (Jedema *et al.*, 2003), comparing favorably with the theoretical $a_{th}^{ph} = 1.2 \times 10^{-4}$.

Spin relaxation in Al depends rather strongly on magnetic fields at low T . CESR measurements (Lubzens

⁸² At some symmetry points $|b|$ may be very small. This occurs in the noble metals which have Fermi states at the Brillouin-zone boundaries, where ΔE is large, but the corresponding λ_{so} is very small due to symmetry.

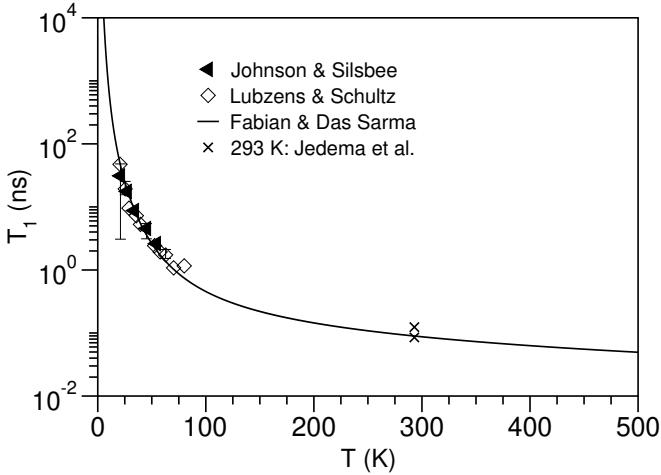


FIG. 16 Measured and calculated τ_s in Al. The low- T measurements are CESR (Lubzens and Schultz, 1976b) and spin injection (Johnson and Silsbee, 1985). Only the phonon contribution is shown, as adapted from Johnson and Silsbee (1985). The solid line is the first-principles calculation, not a fit to the data, (Fabian and Das Sarma, 1998). The data at $T = 293$ K are results from room-temperature spin injection experiments of Jedema *et al.* (2002b, 2003). Adapted from Fabian and Das Sarma, 1999b.

and Schultz, 1976a,b) show that at temperatures below 100 K, $1/\tau_s$ increases linearly with increasing B . A specific sample (Lubzens and Schultz, 1976b) showed a decrease of τ_s from about 20 ns to 1 ns, upon increase in B from 0.05 to 1.4 T. It was proposed that the observed behavior was due to cyclotron motion through spin hot spots (Silsbee and Beuneu, 1983). The reasoning is as follows. Assume that there is considerable spread (anisotropy) $\delta g \approx \Delta g$ of the g factors over the Fermi surface. Such a situation is common in polyvalent metals, whose spin hot spots have anomalously large spin-orbit coupling. In a magnetic field the electron spins precess correspondingly with rates varying by $\delta\Omega_L \approx (\delta g/g)\Omega_L$, where Ω_L is the Larmor frequency. Motional narrowing leads to $1/\tau_s \approx (\delta\Omega_L)^2\tau_c$, where τ_c is the correlation time for the random changes in g . At small magnetic fields $\tau_c = \tau_p$ and $1/\tau_s \sim B^2\tau_p$. Such a quadratic dependence of $1/\tau_s$ on B is a typical motional narrowing case and has been observed at low temperatures in Cu (Lubzens and Schultz, 1976b). As the field increases τ_c becomes the time of flight through spin hot spots, in which case $\tau_c \sim 1/B$. As a result $1/\tau_s$ acquires a component linear in B , in accord with experiment.

In an effort to directly detect phonon-induced spin flips in Al, an interesting experiment was devised (Grimaldi and Fulde, 1996; Lang *et al.*, 1996) using the Zeeman splitting of the energy gap in Al superconducting tunnel junctions. Although the experiment failed, due to overwhelming spin-flip boundary scattering, it showed the direction for future research in studying spin-flip electron-phonon interactions.

D. Spin relaxation in semiconductors

Although sorting out different spin-relaxation mechanisms of conduction electrons in semiconductors is a difficult task, it has generally been observed that the EY mechanism is relevant in small-gap and large-spin-orbit coupling semiconductors, while the DP processes are responsible for spin dephasing in middle-gap materials and at high temperatures. In heavily p-doped samples the BAP mechanism dominates at lower temperatures, while DP at higher. In low-doped systems the DP dominates over the whole temperature range where electron states are extended. Spin relaxation of bound electrons proceeds through the hyperfine interaction. Finally, spin relaxation of holes is due to the EY processes. In bulk III-V or II-VI materials, for holes $\tau_s \approx \tau_p$, since the valence spin and orbital states are completely mixed. However, in two-dimensional systems, where the heavy and light hole states are split, hole spin relaxation is much less effective.

1. Bulk semiconductors

There is a wealth of useful data on τ_s in semiconductors.⁸³

A comprehensive theoretical investigation of spin dephasing in bulk semiconductors (both p- and n-types), applied to GaAs, GaSb, InAs, and InSb, has been carried out by Song and Kim (2002) by using the EY, DP, and BAP mechanisms. The calculation uses analytical formulas like Eq. (66), while explicitly evaluating τ_p for different momentum scattering processes at different control parameters (temperature and density), but only for nondegenerate electron systems (Boltzmann statistics) and zero magnetic field. The main results are as follows: in n-type III-V semiconductors DP dominates at $T \gtrsim 5$ K. At lower T the EY mechanism becomes relevant. This crossover temperature appears to be quite insensitive to the electron density, being between 1 and 5 K in most investigated III-V semiconductors for donor densities greater than 10^{14} cm⁻³ (Song and Kim, 2002). For p-type materials the dominant mechanisms are DP and BAP, with the crossover temperature sensitive to the acceptor density. For example, in p-GaAs at room temperature, the DP mechanism dominates below 10^{18}

⁸³ References for selected semiconductors include the following: p-GaAs: (Aronov *et al.*, 1983; Fishman and Lampel, 1977; Marushchak *et al.*, 1984; Sanada *et al.*, 2002; Seymour *et al.*, 1981; Zerrouati *et al.*, 1988); n-GaAs: See III.D.3; p-Al_xGa_{1-x}As: (Clark *et al.*, 1975; Garbuzov *et al.*, 1971); p-GaSb: (Aronov *et al.*, 1983; Safarov and Titkov, 1980; Sakharov *et al.*, 1981); n-GaSb: (Kauschke *et al.*, 1987); n-InSb: (Chazalviel, 1975); InAs: (Bogess *et al.*, 2000); p-InP: (Gorelenok *et al.*, 1986); n-InP: (Kauschke *et al.*, 1987); n-GaN: (Beschoten *et al.*, 2001; Fanciulli *et al.*, 1993).

cm^{-3} , while at large densities the BAP mechanism dominates. In small-gap InSb the DP mechanism appears to be dominant for all acceptor densities at temperatures above 50 K. The strong disagreement with experiment found at low T (at 5 K, to be specific) points to our still limited theoretical understanding of spin relaxation in semiconductors. The discrepancy likely arises from neglect of HFI effects.

Spin relaxation times as long as 300 ns were recently obtained in bulk, ≈ 100 nm wide, GaAs at low 4.2 K, placed in the proximity of QW's (Dzhioev *et al.*, 2002b, 2001). The samples were low doped ($\approx 10^{14} \text{ cm}^{-3}$ uncompensated donor density), so that optical orientation detected τ_s of electrons bound on donors. At such low donor concentrations the HFI mechanism is responsible for spin relaxation. The unusually large τ_s is attributed to the presence of additional conduction electrons in the structure, coming from the barriers separating the sample and the nearest QW. The hyperfine interaction is then motionally narrowed by the exchange interaction between the donor-bound and conduction electrons. Upon depletion of the conduction electrons from the sample by resonant excitations in the QW, τ_s decreased to 5 ns (Dzhioev *et al.*, 2002b), implying that the effects of the static hyperfine fields on bound-electrons spin precessions are not reduced by motional narrowing.

Spin relaxation of holes in bulk III-V materials is very fast due to a complete mixing of orbital and spin degrees of freedom in the valence band. The EY mechanism predicts that hole τ_s is similar to hole τ_p ; this is a common assumption when considering hole contribution to spin-polarized transport. Hole spin lifetime in undoped GaAs has been measured by optical orientation and time-resolved spectroscopy (Hilton and Tang, 2002). The observed value at room temperature is $\tau_s \approx 110$ fs, consistent with the theoretical assumption.

Spin relaxation of conduction electrons in strained III-V crystals was studied experimentally and theoretically by D'yakonov *et al.* (1986). Spin relaxation under strain is enhanced and becomes anisotropic due to the strain-induced spin splitting of the conduction band, which is linear in k , similarly to the bulk inversion asymmetry in two-dimensional systems (see Sec. III.B.2). It was found that $1/\tau_s \sim \sigma^2$, where σ is the applied stress, and that τ_s is only weakly temperature dependent. Spin relaxation of photoholes in strain crystals has been studied in (D'yakonov and Perel', 1973a), with the conclusion that the hole spin along the strain axis can relax (by the EY processes) on time scales much longer than in unstrained samples, due to the lifting of heavy and light hole degeneracy.

Compared to III-V or II-VI, much less effort has been devoted to investigation of τ_s in bulk Si. The reason is that CESR is thus far the only technique capable of effective detection of spin relaxation in Si. Optical orientation is rather weak (Lampel, 1968) due to the indirect band-gap structure, while robust spin injection in Si is yet to be demonstrated. Spin relaxation in Si is slow due

to the presence of inversion symmetry (the DP mechanism is not applicable) and lack of a nuclear moment for the main Si isotope. Earlier experimental studies (Fehér and Gere, 1959) were concerned with the hyperfine-interaction-dominated spin dephasing in donor states.

A comprehensive experimental study of low-doped Si (P donors were present at the levels $7.5 \times 10^{14} \leq N_d \leq 8 \times 10^{16} \text{ cm}^{-3}$), at temperatures $20 < T < 300$ K, was performed by Lepine (1970). Three distinct temperature regimes were observed:

(a) ($20 < T < 75$ K) Here τ_s decreases with increasing T . The HFI mechanism dominates: electrons are bound to the ground donor states, while thermal excitations to higher states and the exchange interaction with conduction electrons motionally narrows the hyperfine interaction.

(b) ($75 < T < 150$ K) In this temperature range τ_s continues to decrease with increasing T , the effect caused by the spin-orbit interaction in the first excited donor state being motionally narrowed by thermal motion.

(c) ($T > 150$ K) Here $1/\tau_s$ increases with T , in accord with the EY mechanism. The observed room-temperature CESR linewidth is about 8 G, corresponding to the electron spin lifetime of 7 ns.

2. Low-dimensional semiconductor structures

The importance of low-dimensional semiconductor systems (quantum wells, wires, and dots) lies in their great flexibility in manipulating charge and, now, also spin properties of the electronic states. Studies of spin relaxation in those systems are driven not only by the need for fundamental understanding of spin relaxation and decoherence, but also by the goal of finding ways to reduce or otherwise control spin relaxation and coherence in general. For a survey of spin relaxation properties of semiconductor quantum wells, see Sham (1993).

Spin relaxation in semiconductor heterostructures is caused by random magnetic fields originating either from the base material or from the heterostructure itself. All four mechanisms of spin relaxation can be important, depending on the material, doping, and geometry. The difference from the bulk is the localization of the wave function into two, one, or zero dimensions and the appearance of structure-induced random magnetic fields. Of all the mechanisms, the DP and HFI are believed to be most relevant.

The most studied systems are GaAs/AlGaAs QW's. The observed τ_s varies from nanoseconds to picoseconds, depending on the range of control parameters such as temperature, QW width or confinement energy E_1 , carrier concentration, mobility, magnetic field, or bias.⁸⁴

⁸⁴ Here is a list of selected references with useful data on τ_s in GaAs/AlGaAs QW's: confinement energy dependence has been

Spin relaxation has also been investigated in In/GaAs (Cortez *et al.*, 2002; Paillard *et al.*, 2001), in an InAs/GaSb superlattice (Olesberg *et al.*, 2001), in In-GaAs (Guettler *et al.*, 1998), in GaAsSb multiple QW's by (Hall *et al.*, 1999). II-VI QW's (specifically ZnCdSe) were studied by Kikkawa *et al.* (1997), finding $\tau_s \approx 1$ ns, weakly dependent on both mobility and temperature (in the range $5 < T < 270$ K). Electron and hole spin dephasing have also been investigated in dilute magnetic semiconductor QW's doped with Mn ions (Camilleri *et al.*, 2001; Crooker *et al.*, 1997).

Reduction of spin relaxation by inhibiting the BAP electron-hole exchange interaction through spatially separating the two carriers has been demonstrated in δ -doped p-GaAs:Be/AlGaAs (Wagner *et al.*, 1993). The observed τ_s was ≈ 20 ns at $T < 10$ K, which is indeed unusually large. The exchange interaction was also studied at room temperature, observing an increase of τ_s with bias voltage which increases spatial separation between electrons and holes, reducing the BAP effects (Gotoh *et al.*, 2000). In the fractional quantum Hall effect regime it was demonstrated (Kuzma *et al.*, 1998) that nonequilibrium spin polarization in GaAs QW's can survive for tens of μ s. Spin lifetime was also found to be enhanced in GaAs QW's strained by surface acoustic waves (Sogawa *et al.*, 2001). A theoretical study (Kiselev and Kim, 2000) proposed that spin dephasing in 2DEG can be significantly suppressed by constraining the system to finite stripes, several mean free paths wide.

Theoretical studies focusing on spin dephasing in III-V and II-VI systems include those of (Bronold *et al.*, 2002; Krishnamurthy *et al.*, 2003; Lau and Flatté, 2002; Lau *et al.*, 2001; Puller *et al.*, 2003; Wu, 2001; Wu and Kuwata-Gonokami, 2002; Wu and Metiu, 2000). Spin relaxation due to the DP mechanism with bulk inversion asymmetry term in the important case of GaAs/AlGaAs rectangular QW's was investigated by Monte-Carlo simulations (Bournel *et al.*, 2000) at room temperature, including interface roughness scattering. Nice agreement with experiment was found for $\tau_s(E_1)$, where E_1 is the confinement energy. Interface roughness becomes important at large values of E_1 , where scattering increases τ_s (see also Sherman (2003b)).

Spin relaxation and spin coherence of spin-polarized photoexcited electrons and holes in symmetric p- and n-doped and undoped GaAs/AlGaAs quantum wells was investigated using rate equations (Uenoyama and Sham, 1990a,b). It was shown that in these heterostructures

hole spin relaxation proceeds slower than electron-hole recombination. Hole relaxation is found to occur mostly due to acoustic phonon emission. The ratio of the spin-conserving to spin-flip hole relaxation times was found to be 0.46, consistent with the fact that luminescence is polarized even in n-doped quantum wells at times greater than the momentum relaxation time. Similar observations hold for strained bulk GaAs, where hole spin relaxation is also reduced. Spin relaxation of holes in quantum wells was calculated (Bastard and Ferreira, 1992; Ferreira and Bastard, 1991) using the interaction with ionized impurities and s-d exchange in semimagnetic semiconductors. It was shown that size quantization significantly reduces spin relaxation of holes, due to the lifting of heavy and light hole degeneracy. The observed spin lifetimes for holes at low temperatures reached up to 1 ns, while at $T > 50$ K in the same samples τ_s got smaller than 5 ps (Baylac *et al.*, 1995).

Spin dynamics and spin relaxation of excitons in GaAs (Munoz *et al.*, 1995; Vina *et al.*, 2001) and ZnSe (Kalt *et al.*, 2000) were investigated experimentally and theoretically (Maialle *et al.*, 1993; Sham *et al.*, 1998). Coherent spin dynamics in magnetic semiconductors was considered by Linder and Sham (1998).

Spin relaxation in Si heterostructures has been investigated by electron spin resonance in modulation doped Si/SiGe QW's. Very high mobility (about 10^5 cm 2 .V $^{-1}$.s $^{-1}$) samples with $n \approx 3 \times 10^{11}$ cm $^{-2}$ free electrons forming a 2DEG show, at $T = 4.2$ K, T_1 up to 30 μ s (Graeff *et al.*, 1999; Sanderfeld *et al.*, 2000) and T_2 of the order of 100 ns (Graeff *et al.*, 1999), depending on the orientation of B with respect to the QW growth direction. Spin relaxation was attributed to Bychkov-Rashba spin splitting in these asymmetric wells, estimating the corresponding $\hbar\alpha_{BR}$ in Eq. (88) to be around 1×10^{-14} eV.m (Wilamowski and Jantsch, 2002; Wilamowski *et al.*, 2002). Si/Ge heterostructures may have enhanced rates of spin relaxation due to the leakage of the electron wave function to Ge, which is heavier than Si and has greater spin-orbit interaction. Recent studies (Wilamowski and Jantsch, 2004) have confirmed the dominant role of the DP spin relaxation mechanism, leading to the microsecond spin relaxation times. The spin dephasing is argued to be strongly suppressed by cyclotron motion in high-mobility samples (see Sec. III.B.2.a for a brief discussion of the influence of magnetic field on τ_s). Spin-orbit coupling in symmetric Si/SiGe quantum wells has been studied theoretically by Sherman (2003a).

In quantum dots the relevant spin relaxation mechanism is still being debated, as the mechanisms (EY and DP) effective for conduction electrons are ineffective for states localized in QD's (Khaetskii, 2001; Khaetskii and Nazarov, 2000, 2001). It is believed, however, that similar to electrons bound on donors, the dominant mechanism is a HFI process (de Sousa and Das Sarma, 2003a,c; Khaetskii *et al.*, 2002; Merkulov *et al.*, 2002; Semenov and Kim, 2003). Unfortunately, experiments on CdSe QD's (of diameter 22-80 Å) show strong inhomogeneous

studied by Britton *et al.* (1998); Endo *et al.* (2000); Malinowski *et al.* (2000); Ohno *et al.* (2000c, 1999a); Tackeuchi *et al.* (1996); temperature dependence is treated by Adachi *et al.* (2001); Malinowski *et al.* (2000); Ohno *et al.* (2000c, 1999a); Wagner *et al.* (1993); carrier concentration dependence is studied by Sandhu *et al.* (2001); dependence on mobility is examined by Ohno *et al.* (1999a); and dependence on magnetic field in studied by Zhitoimirskii *et al.* (1993).

dephasing ($\tau_s \approx 3$ ns at $B = 0$, while $\tau_s \approx 100$ ps at 4 T) (Gupta *et al.*, 1999), masking the intrinsic spin dephasing processes. Recently a lower bound, limited by the signal-to-noise ratio, on T_1 of 50 μ s has been measured at 20 mK in a one-electron quantum dot defined in 2DEG GaAs/AlGaAs heterostructure by Hanson *et al.* (2003b). The magnetic field of 7.5 T was oriented parallel to the plane heterostructure. While the actual value of T_1 may be orders of magnitude larger, the observed bound suffices for performing elementary quantum gates (see Sec. IV.F).

3. Example: spin relaxation in GaAs

We review recent experimental results on spin relaxation in bulk n-GaAs⁸⁵ and GaAs-based low-dimensional systems.

a. Bulk n-GaAs. The importance of GaAs for spintronics and quantum computing applications has been recently underlined by the discovery of rather long spin relaxation times (of the order of 100 ns) in n-doped samples, as well as by the development of experimental techniques to manipulate spin precession in this semiconductor in a coherent manner (Awschalom, 2001; Oestreich *et al.*, 2002).

Both optical orientation and time-resolved Faraday rotation spectroscopy have been used to measure τ_s in bulk n-GaAs. In the earliest observations of optical spin orientation of electrons, in n-Ga_{0.7}Al_{0.3}As with $N_d \approx 1 \times 10^{16}$ cm⁻³ at 4.2 K, it was found that $\tau_s \approx 1.2$ ns (Ekimov and Safarov, 1971). A much larger spin lifetime was found by optical orientation on n-GaAs (Dzhioev *et al.*, 1997), where for $N_d \approx 1 \times 10^{15}$ cm⁻³ the observed $\tau_s \approx 42$ ns. Faraday rotation studies (Awschalom, 2001; Kikkawa and Awschalom, 1998) found even longer spin lifetimes. At the doping density $N_d = 1 \times 10^{16}$ cm⁻³ of Si donors and $T = 5$ K, the observed $\tau_s \approx 130$ ns at zero magnetic field. At greater and smaller doping densities, spin relaxation time is significantly reduced: for both a nominally undoped sample and for $N_d = 1 \times 10^{18}$ cm⁻³, $\tau_s \approx 0.2$ ns. A comprehensive theoretical investigation of τ_s in bulk n-GaAs is reported by Wu and Ning (2000), and Wu (2001), who solved numerically kinetic equations in the presence of magnetic field. Only the DP mechanism was considered, acting with longitudinal phonon and impurity scattering.

A recent comprehensive study of τ_s based on optical orientation revealed a nice, albeit complex, picture of spin relaxation in bulk n-GaAs over a large range of doping levels (Dzhioev *et al.*, 2002c). Figure 17 summarizes

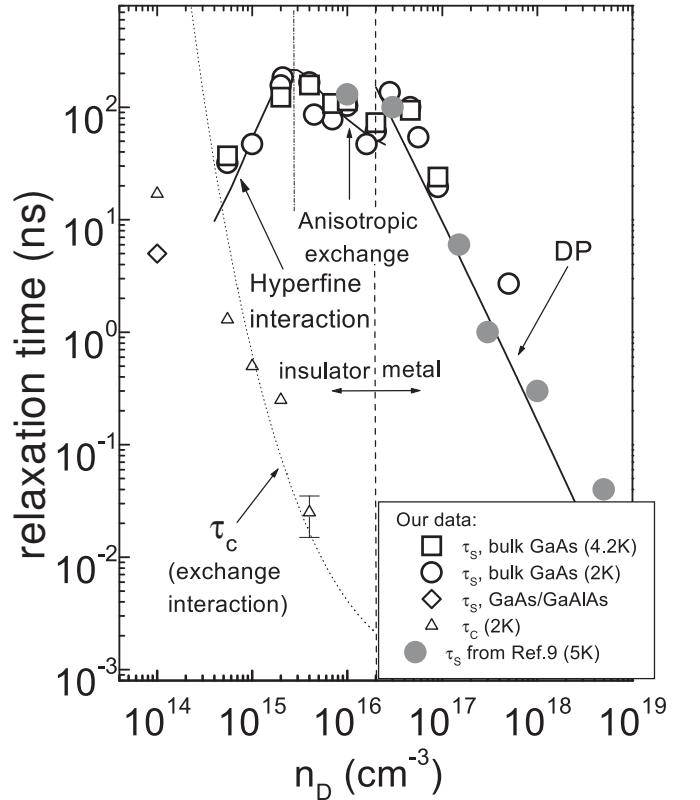


FIG. 17 Spin relaxation time in n-GaAs as a function of donor density N_d (labeled as n_D here) at low temperatures: empty symbols, the optical orientation data of Dzhioev *et al.* (2002a); solid circles are the time-resolved Faraday rotation data of Awschalom (2001); Kikkawa and Awschalom (1998); open triangles, single-spin decoherence times $\tau_{sc} \approx \tau_c$ due to the exchange interaction between electron spins on neighboring donors; solid lines, parameter-free theoretical estimates with labels indicating the dominant spin relaxation mechanisms; dotted line a fit to the experimental data on the exchange correlation time (triangles) τ_c , using a simple model of the exchange coupling between donor states; dashed vertical line, the metal-insulator transition at $N_{dc} = 2 \times 10^{16}$ cm⁻³. From Dzhioev *et al.*, 2002a.

rizes these findings. The spin relaxation time rises with increasing N_d at small doping levels, reaching its first maximum (180 ns) at around 3×10^{15} cm⁻³; τ_s then decreases until $N_d = N_{dc} = 2 \times 10^{16}$ m⁻³, where a sudden increase brings τ_s to another maximum, reaching ≈ 150 ns. At still higher doping levels τ_s decreases strongly with increasing doping.

The above picture is valid at $T \leq 5$ K, where isolated shallow donors are not normally ionized, and the sample is a Mott insulator at small dopings. Conductivity is due to hopping between donor states. Beyond the critical density $N_{dc} \approx 2 \times 10^{16}$ cm⁻³ (the dashed vertical line in Fig. 17) the donor states start to overlap and form an impurity conduction band—electronic states delocalize and the sample becomes metallic. Figure 17 shows that it is the rather narrow window around the metal-to-insulator

⁸⁵ p-GaAs is extensively discussed by Meier and Zakharchenya (Eds.) (1984)

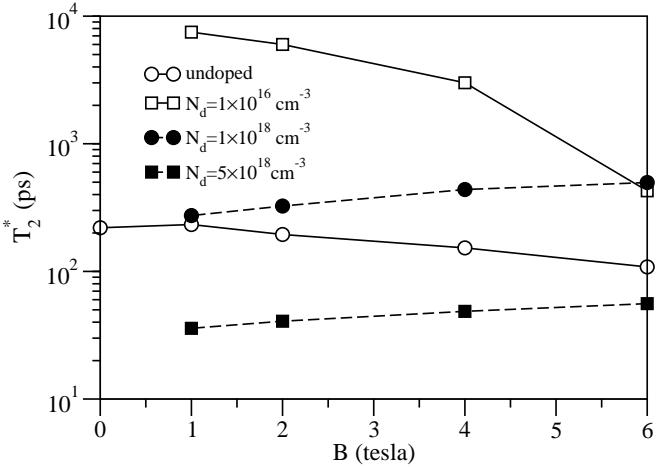


FIG. 18 Measured magnetic-field dependence of the spin dephasing time (here denoted as T_2^* to indicate the likely presence of inhomogeneous broadening; see Sec. III.A.1) for bulk n-GaAs at 5 K. Doping levels, varying from insulating ($N_d < N_{dc} = 2 \times 10^{16} \text{ cm}^{-3}$) to metallic ($N_d > N_{dc}$), are indicated. Adapted from Kikkawa and Awschalom, 1998.

transition where the largest τ_s are found.

At $N_d > N_{dc}$ the DP mechanism dominates. Equation (73) for degenerate electrons explains the observed data rather well. Indeed, considering that $E_F \sim N_d^{2/3}$ and assuming the Brooks-Herring formula for the impurity scattering ($1/\tau_p \sim N_d/E_F^{3/2}$), one obtains $\tau_s \sim 1/N_d^2$, which is observed in Fig. 17. The EY mechanism, Eq. (66), would give $\tau_s \sim N_d^{-4/3}$. The data on the insulating side are consistent with the HFI mechanism: the precession due to local random magnetic fields from the nuclear moments is motionally narrowed by the exchange interaction, which increases with increasing N_d (that is, with increasing overlap between donor states). The theoretical estimates (Dzhioev *et al.*, 2002c) agree well with the data. The behavior of τ_s in the intermediate regime, $3 \times 10^{15} \text{ cm}^{-3} < N_d < N_{dc}$, where τ_s decreases with increasing N_d , was proposed by (Kavokin, 2001) to be due to motional narrowing of the antisymmetric exchange interaction⁸⁶ between bound electrons, with the correlation time τ_c provided by the usual $\mathbf{S}_1 \cdot \mathbf{S}_2$, direct exchange. This new mechanism of spin relaxation, which should be generally present for bound electrons in systems lacking inversion symmetry (such as III-V and II-VI), although still being investigated (Gorkov and Krotkov, 2003; Kavokin, 2002b), appears to give a satisfactory explanation for the experimental data.

In addition to the doping dependence of τ_s , both the temperature and the magnetic field dependences of spin

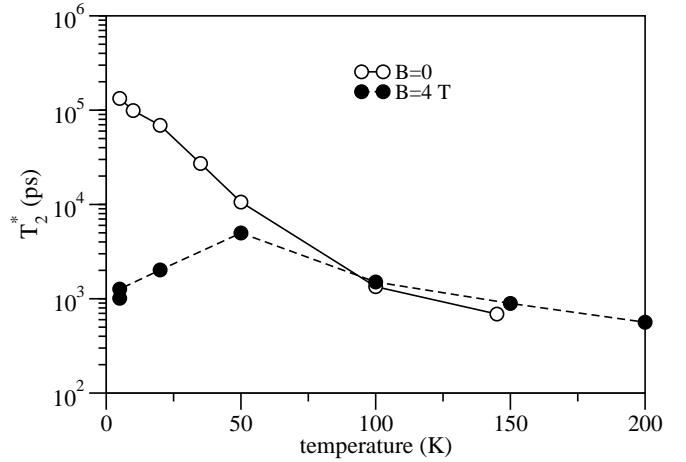


FIG. 19 Measured temperature dependence of the spin dephasing time for bulk n-GaAs doped with $N_d = 1 \times 10^{16} \text{ cm}^{-3}$ Si donors, at $B = 0$ and $B = 4$ T. The sample is insulating at low T and nondegenerate at high T ($T \gtrsim 50$ K, assuming ≈ 4 meV for the donor binding energy), where donors are ionized. Adapted from Kikkawa and Awschalom, 1998.

relaxation in bulk n-GaAs have been studied by Kikkawa and Awschalom (1998). Figure 18 shows $\tau_s(B)$ for samples with varying doping levels at $T = 5$ K. The spin relaxation time increases with B in the metallic regime, the behavior qualitatively consistent with the predictions of the DP mechanism. In contrast, τ_s in the insulating samples decreases with increasing B . Bound electrons are more susceptible to g -factor anisotropies (due to the distribution of electron energies over donor states) and local magnetic field variations (due to the hyperfine interactions). These anisotropies are amplified by increasing B and motionally narrowed by the exchange interaction. It is thus likely that $\tau_s \sim B^{-2}\tau_c(B)$, where the exchange correlation time τ_c depends on B through magnetic orbital effects on the bound electron wave functions (magnetic confinement reduces the extent of the bound orbital, thus reducing the exchange integrals between neighboring donor states). However, no satisfactory quantitative explanation for $\tau_s(B)$ in insulating samples exists.

Figure 19 plots $\tau_s(T)$ for an insulating sample with $N_d = 1 \times 10^{16} \text{ cm}^{-3}$ at $B = 0$ and $B = 4$ T. For the zero-field data the initial decrease of τ_s with B is very rapid, dropping from 130 ns at 5 K to less than 1 ns at 150 K. However, the sample held at $B = 4$ T shows at first a rapid increase with increasing T , and then a decrease at $T \approx 50$ K. The decrease of τ_s with increasing T above 50 K has been found to be consistent with the DP mechanism (Kikkawa and Awschalom, 1998), taking $\tau_s \sim T^{-3}$ in Eq. (74), while extracting the temperature dependence of τ_p from the measurement of mobility. The DP mechanism for conduction electrons was also observed in p-GaAs in the regime of nondegenerate hole densities $N_a \approx 10^{17} \text{ cm}^{-3}$ at temperatures above 100 K (Aronov *et al.*, 1983), after the contribution from the BAP mechanism was subtracted using a the-

⁸⁶ The anisotropic exchange interaction of the Dzyaloshinskii-Moriya form, $\mathbf{S}_1 \times \mathbf{S}_2$ (Dzyaloshinskii, 1958; Moriya, 1960) appears as a result of spin-orbit coupling in semiconductors lacking inversion symmetry.

oretical prediction. From the observed mobility it was found that $\tau_p(T) \sim T^{-0.8}$, so that according to the DP mechanism $\tau_s \sim T^{-2.2}$, which is indeed consistent with the experimental data. The origin of $\tau_s(T)$ below 50 K in Fig. 19 is less obvious. At low T , electrons are localized, so in order to explain the experimental data the theory should include ionization of donors. The increase with increasing T of τ_s at 4 T invokes a picture of motional narrowing in which the correlation time decreases with increasing T much faster than the dispersion of local Larmor frequencies. We do not know of a satisfactory quantitative explanation for these experimental results.⁸⁷ Similar behavior of $\tau_s(T)$ in insulating samples was found in GaN (Beschoten *et al.*, 2001).

The temperature dependence of τ_s for samples with $N_d \gg N_{dc}$ has been reported (Kikkawa and Awschalom, 1998) to be very weak, indicating, for these degenerate electron densities, that $\tau_p(T)$ is only weakly dependent on T . What can be expected for τ_s at room temperature? The answer will certainly depend on N_d . Recent experiments on time-resolved Kerr rotation (Kimel *et al.*, 2001) suggest that $5 \text{ ps} < \tau_s < 10 \text{ ps}$ for undoped GaAs and $15 \text{ ps} < \tau_s < 35 \text{ ps}$ for a heavily doped n-GaAs with $N_d = 2 \times 10^{18} \text{ cm}^{-3}$.

For spintronic applications to make use of the large τ_s observed in bulk n-GaAs one is limited to both very small temperatures and small doping levels. Although this may restrict the design of room-temperature spintronic devices, such a regime seems acceptable for spin-based quantum computing (see Sec. IV.F), where one is interested in the spin lifetime of single (or a few) electrons, bound to impurities or confined to quantum dots. How close is τ_s to the individual spin lifetime τ_{sc} ? There is no clear answer yet. Ensemble spin dephasing seen for insulating GaAs samples appears to be due to motional narrowing of the hyperfine interaction. The randomizing processes are spin flips due to the direct exchange, lead-

ing to the correlation time τ_c , which can be taken as a measure for the lifetime τ_{sc} of the individual spins. Extracting this lifetime from the experiment is not easy, but the obvious trend is the smaller the τ_c , the larger the τ_s . For a specific model of spin relaxation in bound electron states τ_c was extracted experimentally by Dzhioev *et al.* (2002c) by detecting the changes in the spin polarization due to longitudinal magnetic fields. The result is shown in Fig. 17. The two times, τ_c and τ_s differ by orders of magnitude. For the doping levels where τ_s is greater than 100 ns, τ_c is smaller than 0.1 ns. Unfortunately, the useful time for spin quantum computing would be extracted in the limit of very small dopings, where the data are still sparse. For an informal recent review of τ_s in n-GaAs, see Kavokin (2002a).

Closely related to spin relaxation is spin diffusion. Hägele *et al.* (1998) observed the transport of a spin population—longitudinal spin drift—in i-GaAs over a length scale greater than 4 μm in electric fields up to 6 kV/cm and at low temperatures. This was followed by a remarkable result of Kikkawa and Awschalom (1999), the observation of the drift of precessing electron spins—transverse spin drift—in GaAs with $N_d = 1 \times 10^{16} \text{ cm}^{-3}$, over 100 μm in moderate electric fields (tens of V/cm) at $T = 1.6 \text{ K}$, setting the length scale for the spin dephasing. By directly analyzing the spreading and drifting of the electron spin packets in time, Kikkawa and Awschalom obtained the spin diffusion (responsible for spreading) and electronic diffusion (drift by electric field) coefficients. It was found that the former is about 20 times as large as the latter. These results are difficult to interpret, since the sample is just below the metal-to-insulator transition, where charge is transported via hopping, but they suggest that spin diffusion is strongly enhanced through the exchange interaction. Investigations of this type in even smaller doping limits may prove important for understanding single-spin coherence.

⁸⁷ There is a discrepancy in the data presented in Figs. 18 and 19. Take the $N_d = 1 \times 10^{16} \text{ cm}^{-3}$ sample. While Fig. 18 reports $\tau_s \approx 3 \text{ ns}$ at 5 K and 4 T, τ_s is only about 1 ns in Fig. 19. The reason for this difference (Kikkawa, 2003) turns out to be electronically-induced nuclear polarization (Kikkawa and Awschalom, 2000). At low temperatures and large magnetic fields, nuclear polarization develops via the Overhauser effect inhomogeneously throughout the electron spin excitation region. The inhomogeneous magnetic field due to polarized nuclei causes inhomogeneous broadening of the electronic τ_s . The measured spin dephasing time is indeed T_2^* , rather than the intrinsic T_2 . Furthermore, since nuclear polarization typically takes minutes to develop, the measured T_2^* depends on the measurement “history.” This is the reason why two different measurements, reported in Figs. 18 and 19, show different T_2^* under otherwise equivalent conditions. The nuclear polarization effect is also part of the reason why the $T_2(T)$ at 4 T sharply deviates from that at zero field at small T . The technique should give consistent results at small fields and large temperatures, as well as in heavily doped samples where the nuclear fields are motionally narrowed by the itinerant nature of electrons.

b. *GaAs-based quantum wells.* We discuss selected experimental results on spin relaxation in GaAs/Al_xGa_{1-x}As QW’s, presenting the temperature and confinement energy dependence of τ_s .

Figure 20 plots the temperature dependence of $1/\tau_s$ in the interval of $90 < T < 300 \text{ K}$ for QW’s of width L ranging from 6 to 20 nm (Malinowski *et al.*, 2000). The wells, with $x = 0.35$ and orientation along [001], were grown on a single wafer to minimize sample-to-sample variations when comparing different wells. The reported interface roughness was less than the exciton Bohr radius of 13 nm. In these structures the excitonic effects dominate at $T < 50 \text{ K}$ (with the reported $\tau_s \approx 50 \text{ ps}$), while the exciton ionization is complete roughly at $T > 90 \text{ K}$, so the data presented are for free electrons. Spin relaxation was studied using pump-probe optical orientation spectroscopy with a 2 ps time resolution and the typical excitation intensity/pulse of 10^{10} cm^{-2} .

As Fig. 20 shows, τ_s depends rather weakly on T for the

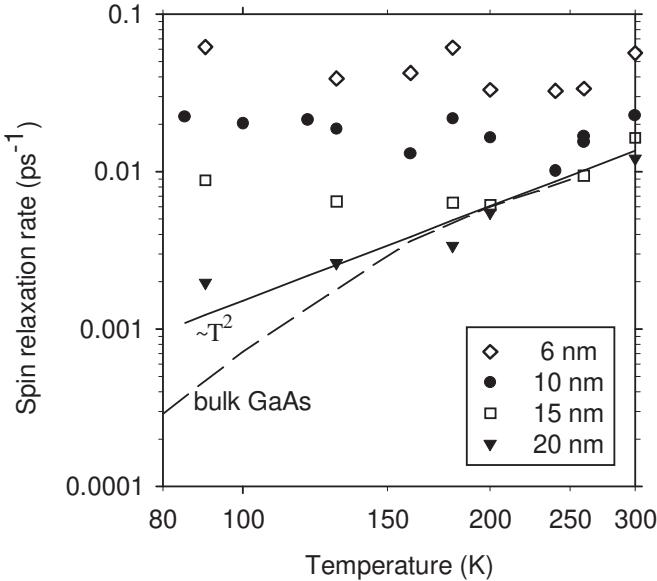


FIG. 20 Measured temperature dependence of the conduction-electron-spin relaxation rate $1/\tau_s$ in GaAs/AlGaAs QW's of varying widths: the dashed curve, data for a low-doped ($N_a = 4 \times 10^{16} \text{ cm}^{-3}$) bulk p-GaAs (Meier and Zakharchenya (Eds.), 1984); solid line, the $\tau_s \sim T^2$ dependence. From Malinowski *et al.*, 2000.

narrow wells with $L < 10 \text{ nm}$. For the well with $L = 15 \text{ nm}$, after being approximately constant (or somewhat decreasing) as T increases to about 200 K, τ_s increases with increasing T at greater temperatures. The increase is consistent with the $1/\tau_s \sim T^2$ behavior. The thickest well increases with the same power law, $1/\tau_s \sim T^2$, over the whole temperature range. In order to make a reliable comparison with theoretical predictions (the expected mechanism is that of DP in two-dimensional systems), one needs to know the behavior of $\tau_p(T)$. The DP mechanism predicts, for the nondegenerate electron densities employed in the experiment, that $1/\tau_s \sim T^3 \tau_p$ [see Eq. (74)] in the bulk and wide QD's, the condition being that thermal energy is greater than the subband separation), and $1/\tau_s \sim TE_1^2 \tau_p$ from Eq. (79) for the bulk inversion asymmetry after making thermal averaging ($E_{\mathbf{k}} \rightarrow k_B T$), when one realizes that confinement energy E_1 is $\sim \langle k_n^2 \rangle$. When one assumes that momentum relaxation in these elevated temperatures is due to scattering by phonons, τ_p should be similar in bulk and low-dimensional structures. From the observed high-temperature bulk $\tau_s(T)$ (at low temperatures τ_s is affected by BAP processes) one can estimate $\tau_p \sim 1/T$, which is consistent with the constant τ_s for the narrow wells, and with the quadratic dependence for the wide wells. At low T , in addition to the BAP mechanism, τ_s will deviate from that in the bulk due to impurity scattering. The EY and BAP mechanisms were found not to be relevant to the observed data (Malinowski *et al.*, 2000).

Figure 21 shows the dependence of $1/\tau_s$ on the experi-

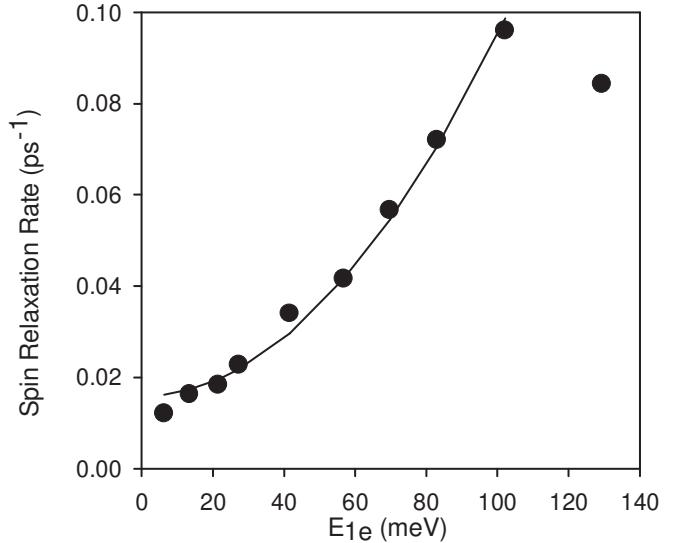


FIG. 21 Measured room-temperature dependence of $1/\tau_s$ on the confinement energy E_1 for GaAs/AlGaAs QW's. The solid line is a quadratic fit, showing behavior consistent with the DP mechanism. From Malinowski *et al.*, 2000.

mentally determined confinement energy E_1 for a variety of QW's on the same wafer (Malinowski *et al.*, 2000). The data are at room temperature. The spin relaxation time varies from somewhat less than 100 ps for wide QW's, approximating the bulk data (cf. Kimel *et al.* (2001) where $15 \text{ ps} < \tau_s < 35 \text{ ps}$ was found for a heavily doped n-GaAs), to about 10 ps in most confined structures. The downturn for the highest- E_1 well (of width 3 nm) is most likely due to the increased importance of interface roughness at such small widths (Malinowski *et al.*, 2000). Confinement strongly enhances spin relaxation. This is consistent with the DP mechanism for two-dimensional systems, in which the spin precession about the intrinsic magnetic fields (here induced by bulk inversion asymmetry) increases as E_1^2 with increasing confinement. The observed data in Fig. 21 are consistent with the theoretical prediction.

Similarly to bulk GaAs, spin relaxation in GaAs QW's was found to be reduced at carrier concentrations close to the metal-to-insulator transition ($n \approx 5 \times 10^{10} \text{ cm}^{-2}$) (Sandhu *et al.*, 2001).

IV. SPINTRONIC DEVICES AND APPLICATIONS

In this section we focus primarily on the physical principles and materials issues for various device schemes, which, while not yet commercially viable, are likely to influence future spintronic research and possible applications.

A. Spin-polarized transport

1. F/I/S tunneling

Experiments reviewed by Tedrow and Meservey (1994) in ferromagnet/insulator/superconductor (F/I/S) junctions have established a sensitive technique for measuring the spin polarization P of magnetic thin films and, at the same time, has demonstrated that the current will remain spin-polarized after tunneling through an insulator. These experiments also stimulated more recent imaging techniques based on the spin-polarized STM (see Johnson and Clarke (1990); Wiesendanger *et al.* (1990); and a review, Wiesendanger (1998)) with the ultimate goal of imaging spin configurations down to the atomic level.

The degree of spin polarization is important for many applications such as determining the magnitude of tunneling magnetoresistance (TMR) in magnetic tunnel junctions (MTJ) [recall Eq. (2)]. Different probes for spin polarization generally can measure significantly different values even in experiments performed on the same homogeneous sample. In an actual MTJ, measured polarization is *not* an intrinsic property of the F region and could depend on interfacial properties and the choice of insulating barrier. Challenges in quantifying P , discussed here in the context of F/I/S tunneling, even when F is a simple ferromagnetic metal, should serve as a caution for studies of novel, more exotic, spintronic materials.

F/I/S tunneling conductance is shown in Fig. 22, where for simplicity we assume that the spin-orbit and spin-flip scattering (see Sec. III.C) can be neglected, a good approximation for $\text{Al}_2\text{O}_3/\text{Al}$ (Tedrow and Meservey, 1971a, 1994), a common choice for I/S regions. For each spin the normalized BCS density of states is $\tilde{\mathcal{N}}_S(E) = \text{Re}(|E|/2\sqrt{E^2 - \Delta^2})$, where E is the quasiparticle excitation energy and Δ the superconducting gap.⁸⁸ The BCS density of states is split in a magnetic field H , applied parallel to the interface, due to a shift in quasiparticle energy $E \rightarrow E \pm \mu_B H$, for \uparrow (\downarrow) spin parallel (antiparallel) to the field, where μ_B is the Bohr magneton. The tunneling conductance is normalized with respect to its normal state value—for an F/I/N junction, $G(V) \equiv (dI/dV)_S/(dI/dV)_N = G_{\uparrow}(V) + G_{\downarrow}(V)$, where V is the applied bias. This conductance can be expressed by generalizing analysis of Giaever and Megerle (1961) as

$$G(V) = \int_{-\infty}^{\infty} \frac{1+P}{2} \frac{\tilde{\mathcal{N}}_S(E + \mu H)\beta dE}{4 \cosh^2[\beta(E + qV)/2]} + \int_{-\infty}^{\infty} \frac{1-P}{2} \frac{\tilde{\mathcal{N}}_S(E - \mu H)\beta dE}{4 \cosh^2[\beta(E + qV)/2]}. \quad (99)$$

Here $\beta = 1/k_B T$, k_B is the Boltzmann constant, T is the temperature, and q is the proton charge. The factors

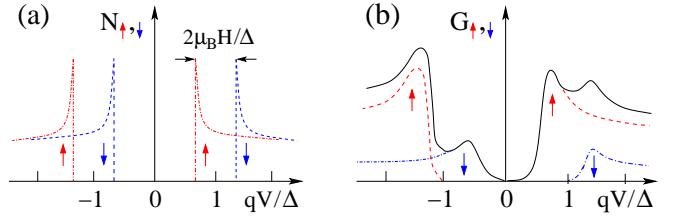


FIG. 22 Ferromagnet/insulator/superconductor tunneling in an applied magnetic field: (a) Zeeman splitting of the BCS density of states as a function of applied bias; (b) normalized spin-resolved conductance (dashed lines) and the total conductance (solid line) at finite temperature.

$(1 \pm P)/2$ represent the difference in tunneling probability between for \uparrow and \downarrow electrons. While a rigorous determination of P , in terms of materials parameters, would require a full calculation of spin-dependent tunneling, including the appropriate boundary conditions and a detailed understanding of the interface properties, it is customary to make some simplifications. Usually P can be identified as (Maekawa *et al.*, 2002; Worledge and Geballe, 2000a)

$$P \rightarrow P_G = (G_{N\uparrow} - G_{N\downarrow})/(G_{N\uparrow} + G_{N\downarrow}), \quad (100)$$

the spin polarization of the normal-state conductance (proportional to the weighted average of the density of states in F and S and the square of the tunneling matrix element), where \uparrow is the electron spin with the magnetic moment parallel to the applied field (majority electrons in F). With the further simplification of spin-independent and constant tunneling matrix element (Tedrow and Meservey, 1971b, 1994), Eq. (100) can be expressed as

$$P \rightarrow P_N = (\mathcal{N}_{F\uparrow} - \mathcal{N}_{F\downarrow})/(\mathcal{N}_{F\uparrow} + \mathcal{N}_{F\downarrow}), \quad (101)$$

the spin polarization of the tunneling density of states in the F region at the Fermi level.

Spin polarization P of the F electrode can be deduced (Tedrow and Meservey, 1994) from the asymmetry of the conductance amplitudes at the four peaks in Fig. 22 (b) [for $P=0$, $G(V)=G(-V)$]. In $\text{CrO}_2/\text{I/S}$ tunnel junctions, nearly complete spin polarization $P_G > 0.9$ was measured (Parker *et al.*, 2002). Only two of the four peaks sketched in Fig. 22, have been observed, indicating no features due to the minority spin up to $H=2.5$ T. Parkin *et al.* (2004) have shown that by replacing an aluminum oxide (a typical choice for an insulating region) with magnesium oxide, one can significantly increase the spin polarization in F/I/S junctions. Correspondingly, extraordinarily large values of TMR (> 200% at room temperature) can be achieved even with conventional ferromagnetic CoFe) electrodes.

The assumption of spin-conserving tunneling can be generalized (Monsma and Parkin, 2000a,b; Tedrow and Meservey, 1994; Worledge and Geballe, 2000a) to extract P in the presence of spin-orbit and spin-flip scattering.

⁸⁸ Here we focus on a conventional *s*-wave superconductor with no angular dependence in Δ .

Theoretical analyses (Bruno and Schwartz, 1973; Fulde, 1973; Maki, 1964) using many-body techniques show that the spin-orbit scattering would smear the Zeeman-split density of states, eventually merging the four peaks into two, while the magnetic impurities (Abrikosov and Gorkov, 1960) act as pair breakers and reduce the value of Δ . Neglecting the spin-orbit scattering was shown to lead to the extraction of higher P values (Monsma and Parkin, 2000a; Tedrow and Meservey, 1994).

With a few exceptions (Worledge and Geballe, 2000b), F/I/S conductance measurements (Tedrow and Meservey, 1994) have revealed positive P —the dominant contribution of majority spin electrons for different ferromagnetic films (for example, in Fe, Ni, Co and Gd). However, electronic structure calculations typically give that $N_{F\uparrow} < N_{F\downarrow}$ and $P_N < 0$ [for Ni and Co $N_{F\uparrow}/N_{F\downarrow} \sim 1/10$ (Butler *et al.*, 2001)]. Early theoretical work addressed this apparent difference,⁸⁹ and efforts to understand precisely what is being experimentally measured have continued.

Stearns (1977) suggested that only itinerant, freelike electrons will contribute to tunneling, while nearly localized electrons, with a large effective mass, contribute to the total density of states but not to $G(V)$ [see also Hertz and Aoi (1973) and, for spin-unpolarized tunneling, Gadzuk (1969)]. From the assumed parabolic dispersion of the spin subbands with fixed spin splitting, Stearns related the measured polarization to the magnetic moment, giving positive $P \rightarrow P_k = (k_{F\uparrow} - k_{F\downarrow})/(k_{F\uparrow} + k_{F\downarrow})$, the spin polarization of the projections of Fermi wave vectors perpendicular to the interface. Similar arguments, for inequivalent density-of-states contributions to $G(V)$, were generalized to more complex electronic structure. Mazin (1999) showed the importance of the tunneling matrix elements which have different Fermi velocities for different bands [see also (Yusof *et al.*, 1998), in the context of tunneling in a high temperature superconductor (HTSC)]. Consequently, P_G could even have an opposite sign from P_N —which, for example, would be measured by spin-resolved photoemission.

Good agreement between tunneling data and electronic structure calculations was illustrated by the example of Ni_xFe_{1-x} (Nadgorny *et al.*, 2000), showing, however, that P is not directly related to the magnetic moment (Meservey *et al.*, 1976). The difference between bulk and the surface densities of states of the ferromagnet (probed in tunneling measurements) (Oleinik *et al.*, 2000), the choice of tunneling barrier (De Teresa *et al.*, 1999), and details of the interfacial properties, which can change over time (Monsma and Parkin, 2000b), have all been shown to affect the measured P directly.

Tedrow-Meservey technique is also considered as a probe to detect spin injection in Si, where optical methods, due to the indirect gap, would be ineffective. F/I/S

tunneling was also studied using amorphous Si (a-Si) and Ge (a-Ge) as a barrier. While with a-Si some spin polarization was detected (Meservey *et al.*, 1982) no spin-polarized tunneling was observed using a-Ge barrier (Gibson and Meservey, 1985), in contrast to the first reports of TMR (Jullierè, 1975).

Spin-dependent tunneling was also studied using a HTSC electrode as a detector of spin polarization (Chen *et al.*, 2001; Vas'ko *et al.*, 1998). While this can significantly extend the temperature range in the tunneling experiments, a lack of understanding of HTSC's makes such structures more a test ground for fundamental physics than a quantitative tool for quantitatively determining P . There are also several important differences between studies using HTSC's and conventional low-temperature superconductors. The superconducting pairing symmetry no longer yields an isotropic energy gap, and even for the BCS-like picture the density of states should be accordingly modified. A sign change of the pair potential can result in $G(V=0) > 0$ for $T \rightarrow 0$ even for a strong tunneling barrier and give rise to a zero-bias conductance peak (Hu, 1994; Tanaka and Kashiwaya, 1995; Wei *et al.*, 1998). This is explained by the two-particle process of Andreev reflection (discussed further in Sec. IV.A.3), which, in addition to the usual quasiparticle tunneling, contributes to the I-V characteristics of a F/I/S junction (Hu and Yan, 1999; Kashiwaya *et al.*, 1999; Zhu *et al.*, 1999; Žutić and Valls, 1999, 2000) [a simpler N/I/S case is reviewed by Hu (1998) and Kashiwaya and Tanaka (2000)]. The suppression of a zero-bias conductance peak, measured by an STM, was recently used to detect spin injection into a HTSC (Ngai *et al.*, 2004).

2. F/I/F tunneling

In the preface to a now classic reference on spin-unpolarized tunneling in solids, Duke (1969) concludes that (with only a few exceptions) the study of tunneling is an art and not a science. Perhaps this is also an apt description for the present state of experiment on spin-polarized tunneling between two ferromagnetic regions. Even for MTJ's with standard ferromagnetic metals, the bias and the temperature dependence of the TMR, as identification of the relevant spin polarization remain to be fully understood. In a brief review of current findings we intend to identify questions that could arise as new materials for MTJ's are being considered.

A resurgence in interest in the study of MTJ's, following a hiatus after the early work by Jullierè (1975); Maekawa and Gäfvert (1982), was spurred by the observation of large room-temperature TMR (Miyazaki and Tezuka, 1995; Moodera *et al.*, 1995). This discovery has opened the possibility of using MTJ's for fundamental studies of surface magnetism and room-temperature spin polarization in various ferromagnetic electrodes as well as to suggesting applications such as highly sensitive

⁸⁹ For a list of references see Tedrow and Meservey (1973, 1994).

magnetic-field sensors, magnetic read heads, and non-volatile magnetic memory applications.

It is instructive to notice the similarity between the schematic geometry and the direction of current flow in an MTJ and that in CPP GMR (recall Figs. 2 and 3), which only differ in the middle layer being an insulator and a metal, respectively. By considering the limit of ballistic transport in CPP GMR⁹⁰ it is possible to give a unified picture of both TMR and CPP GMR by varying the strength of the hopping integrals (Mathon and Umerski, 1997) in a tight-binding representation.

Jullière (1975) modified Eq. (100) in the limit $V \rightarrow 0$, $T \rightarrow 0$, and applied it to study F/I/F tunneling. The two F regions are treated as uncoupled with the spin-conserving tunneling across the barrier. This effectively leads to the two-current model proposed by Mott (1936a) and also applied to the CPP GMR geometries (Gijs and Bauer, 1997; Valet and Fert, 1993). The values for P extracted from F/I/S measurements are in a good agreement with the observed TMR values (typically positive, as expected from $P_{1,2} > 0$). However, Jullière's formula⁹¹ does not provide an explicit TMR dependence on bias and temperature.

Jullière's result can be obtained as a limiting case from a more general Kubo/Landauer approach (Mathon and Umerski, 1999) with the assumption that the component of the wave vector parallel to the interface \mathbf{k}_{\parallel} is not conserved (incoherent tunneling). Such a loss of coherence is good approximation for simply capturing the effects of disorder for amorphous Al₂O₃, a common choice for the I region with metallic ferromagnets. Despite its simplicity, the Jullière's model for the TMR has continued to be used for interpreting the spin polarization in various MTJ's. Recent examples include F regions made of manganite perovskites displaying colossal magnetoresistance (CMR) (Bowen *et al.*, 2003) (suggesting $P_N > 0.95$); magnetite (Fe₃O₄) (Hu and Suzuki, 2002) (with $P < 0$ and TMR < 0); III-V ferromagnetic semiconductors (Chun *et al.*, 2002); a nonmagnetic semiconductor used as a tunneling barrier (Kreuzer *et al.*, 2002); Co/carbon nanotube/Co MTJ (Tsukagoshi *et al.*, 1999); and resonant tunneling in F/I/N/F junctions (Yuasa *et al.*, 2002).

For novel materials, in which the electronic structure calculations and an understanding of the interfacial properties are not available, Jullière's formula still provides useful insights. A quantitative understanding of MTJ's challenges similar to those discussed for F/I/S tunneling, including determining precisely which spin polarization is relevant and the related issue of reconciling the (typically positive) sign of the observed TMR with the electronic structure (Bratkovsky, 1997; LeClair *et al.*, 2002; Ma-

cLaren *et al.*, 1997; Mathon and Umerski, 1997; Oleinik *et al.*, 2000; Tsymbal and Pettifor, 1997).

In an approach complementary to Jullière's, Slonczewski (1989) considered F/I/F as a single quantum-mechanical system in a free-electron picture. When matching the two-component wave functions at interfaces, coherent tunneling was assumed, with conserved \mathbf{k}_{\parallel} , relevant to epitaxially grown MTJ's (Mathon and Umerski, 2001) and the I region was modeled by a square barrier.⁹² The resulting TMR can be expressed as in Eq. (2) but with the redefined polarization

$$P \rightarrow P_k(\kappa^2 - k_{F\uparrow}k_{F\downarrow})/(\kappa^2 + k_{F\uparrow}k_{F\downarrow}), \quad (102)$$

where P_k , as defined by Stearns (1977), is also P_N (in a free-electron picture) and $i\kappa$ is the usual imaginary wave vector through a square barrier. Through the dependence of κ on V the resulting polarization in Slonczewski's model can change sign. A study of a similar geometry using a Boltzmann-like approach shows (Chui, 1997) that the spin splitting of electrochemical potentials persists in the F region all the way to the F/I interface, implying $\kappa_{\uparrow} \neq \kappa_{\downarrow}$ and an additional voltage dependence of the TMR. Variation of the density of states [inferred from the spin-resolved photoemission data (Park *et al.*, 1998a,b)] within the range of applied bias in MTJ's of Co/SrTiO₃/La_{0.7}Sr_{0.3}MnO₃ (Co/STO/LSMO) (De Teresa *et al.*, 1999), together with Jullière's model, was used to explain the large negative TMR (-50% at 5 K), which would even change sign for positive bias (raising the Co Fermi level above the corresponding one of LSMO). The bias dependence of the TMR was also attributed to the density of states by extending the model of a trapezoidal tunneling barrier (Brinkman *et al.*, 1970) to the spin-polarized case (Xiang *et al.*, 2002).

The decay of TMR with temperature can be attributed to several causes. Early theoretical work on N/I/N tunneling (Anderson, 1966; Appelbaum, 1966) [for a detailed discussion and a review of related experimental results see Duke (1969)] showed that the presence of magnetic impurities in the tunneling barrier produces temperature dependent conductance—referred to as zero-bias anomalies. These findings, which considered both spin-dependent and spin-flip scattering, were applied to fit the decay of the TMR with temperature (Inoue and Makeawa, 1999; Jansen and Moodera, 2000; Miyazaki, 2002). Hot electrons localized at F/I interfaces were predicted, to create magnons, or collective spin excitations, near the F/I interfaces, and suppress the TMR (Zhang *et al.*, 1997). Magnons were observed (Tsui *et al.*, 1971) in an antiferromagnetic (AFM) NiO barrier in single crystal Ni/NiO/Pb tunnel junctions and were suggested

⁹⁰ Related applications are usually in a diffusive regime.

⁹¹ For its limitations and extensions see comprehensive reviews by Moodera and Mathon (1999); Moodera *et al.* (1999).

⁹² A formally analogous problem was considered by Griffin and Demers (1971) in an N/I/S system where the two-component wave functions represented electron-like and hole-like quasiparticles rather than the two spin projections; see Sec. IV.A.3.

(Moodera *et al.*, 1995) as the cause of decreasing TMR with T by spin-flip scattering. Using an s - d exchange (between itinerant s and nearly localized d electrons) Hamiltonian, it was shown (Zhang *et al.*, 1997) that, at $V \rightarrow 0$, $G(T) - G(0) \propto T \ln T$, for both $\uparrow\uparrow$ and $\uparrow\downarrow$ orientations. A different temperature dependence of TMR was suggested by Moodera *et al.* (1998). It was related to the decrease of the surface magnetization (Pierce and Celotta, 1984; Pierce *et al.*, 1982) $M(T)/M(0) \propto T^{3/2}$. Such a temperature dependence [known as the Bloch's law and reviewed by Krey (2004)], attributed to magnons, was also obtained for TMR (MacDonald *et al.*, 1998). An additional decrease of TMR with T was expected due to the spin-independent part of $G(T)$ (Shang *et al.*, 1998), seen also in N/I/N junctions.

Systematic studies of MTJ's containing a semiconductor (Sm) region (used as a tunneling barrier and/or an F electrode) have begun only recently.⁹³ To improve the performance of MTJ's it is desirable to reduce the junction resistance. A smaller RC constant would allow faster switching times in MRAM (for a detailed discussion see De Boeck *et al.* (2002)). Correspondingly, using a semiconducting barrier could prove an alternative strategy for difficult fabrication of ultrathin (< 1 nm) oxide barriers (Rippard *et al.*, 2002). Some F/Sm/F MTJ's have been grown epitaxially, and the amplitude of TMR can be studied as function of the crystallographic orientation of a F/Sm interface. For an epitaxially grown Fe/ZnSe/Fe MTJ electronic structure calculations have predicted (MacLaren *et al.*, 1999) large TMR (up to ~ 1000 %), increasing with ZnSe thickness. However, the observed TMR in Fe/ZnSe/Fe_{0.85}C_{0.15} was limited below 50 K, reaching 15 % at 10 K for junctions of higher resistance and lower defect density⁹⁴ (Gustavsson *et al.*, 2001). Results on ZnS, another II-VI semiconductor, demonstrated a TMR of ~ 5 % at room temperature (Guth *et al.*, 2001).

There is also a possibility of using all-semiconductor F/Sm/F single-crystalline MTJ's where F is a ferromagnetic semiconductor. These would simplify integration with the existing conventional semiconductor-based electronics and allow flexibility of various doping profiles and fabrication of quantum structures, as compared to the conventional all-metal MTJ's. Large TMR (> 70 % at 8 K), shown in Fig. 23, has been measured in an epitaxially grown (Ga,Mn)As/AlAs//(Ga,Mn)As junction (Tanaka and Higo, 2001). The results are consistent with

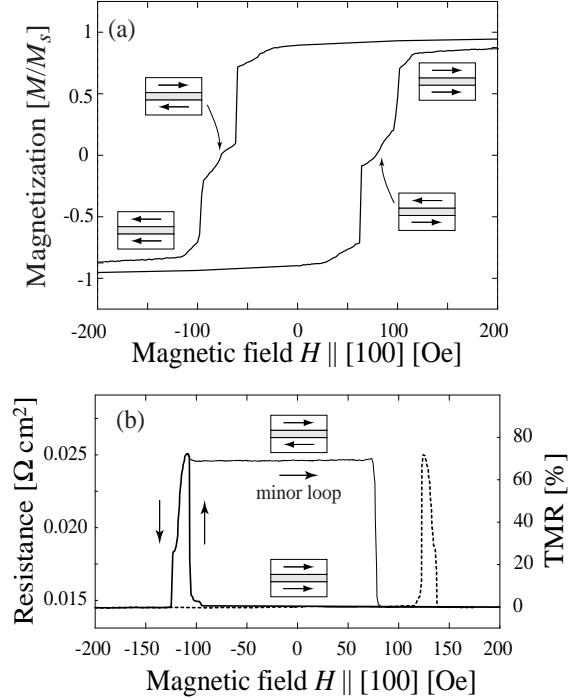


FIG. 23 All-semiconductor magnetic tunnel junction: (a) magnetization of $\text{Ga}_{1-x}\text{Mn}_x\text{As}$ ($x=4.0\%, 50$ nm)/AlAs (3 nm)/ $\text{Ga}_{1-x}\text{Mn}_x\text{As}$ ($x=3.3\%, 50$ nm) trilayer measured by a SQUID at 8 K. The sample size is $3 \times 3 \text{ mm}^2$. Magnetization shown is normalized with respect to the saturation value M_s . (b) TMR curves of a $\text{Ga}_{1-x}\text{Mn}_x\text{As}$ ($x=4.0\%, 50$ nm)/AlAs (1.6 nm)/ $\text{Ga}_{1-x}\text{Mn}_x\text{As}$ ($x=3.3\%, 50$ nm) tunnel junction of $200 \mu\text{m}$ in diameter. Bold solid curve, sweep of the magnetic field from positive to negative; dashed curve, sweep from negative to positive; thin solid curve, a minor loop. From Tanaka and Higo, 2001.

the k_{\parallel} being conserved in the tunneling process (Mathon and Umerski, 1997), with the decrease of TMR with T expected from the spin-wave excitations (MacDonald *et al.*, 1998; Shang *et al.*, 1998), discussed above. TMR is non-monotonic with thickness in AlAs (with the peak at ~ 1.5 nm). For a given AlAs thickness, double MTJ's were also shown to give similar TMR values and were used to determine electrically the spin injection in GaAs QW (Mattana *et al.*, 2003). However, a room-temperature effect remains to be demonstrated as the available well-characterized ferromagnetic semiconductors do not have as high a ferromagnetic transition temperature.

A lower barrier in F/Sm/F MTJ's can have important implications in determining the actual values of TMR. The standard four-probe technique for measuring I and V has been known to give spurious values when the resistance of the F electrodes is non-negligible to the junction resistance. The tunneling current in that regime has been shown to be highly nonuniform⁹⁵ and the measured ap-

⁹³ The early F/Ge/F results (Jullierè, 1975) were not reproduced and other metallic structures involving Si, Ge, GaAs, and GaN as a barrier have shown either no (Boeve *et al.*, 2001; Gibson and Meservey, 1985; Loraine *et al.*, 2000) or only a small (Jia *et al.*, 1996; Kreuzer *et al.*, 2002; Meservey *et al.*, 1982) spin-dependent signal.

⁹⁴ Interface defects could diminish measured TMR. We recall (see Sec. II.D.3) that at a ZnMnSe/AlGaAs interface they limit the spin injection efficiency (Stroud *et al.*, 2002) and from Eq. (32) infer a reduced spin-valve effect.

⁹⁵ Nonuniform tunneling current has been studied in nonmagnetic

parent resistance $R_m = V/I$ (different from the actual junction resistance R_J) can even attain negative values (Moodera *et al.*, 1996; Pederson and Vernon, Jr., 1967). The important implications for MTJ are the possibility of large overestimates in the TMR amplitude (Moodera and Mathon, 1999) and a desirable hysteresis effect—at $H=0$ the two values of resistance can be used for various nonvolatile applications (Moodera *et al.*, 1996).

A detailed understanding of MTJ's will also require knowing the influence of the interface and surface roughness (Itoh *et al.*, 1999). Even in the spin-unpolarized case it is known that the full quantum-mechanical approach (Tešanović *et al.*, 1986) can lead to qualitatively different results from the usual quasiclassical picture and from averaging out the spatial information on the length scale of the inverse Fermi wave vector.

A comprehensive review of tunneling phenomena and magnetoresistance in granular materials, ferromagnetic single-electron transistors, and double tunnel junctions is given by Maekawa *et al.* (2002). A theoretical study of F/I/F junctions, in which the I region is a quantum dot, shows the importance of Coulomb interactions, which could lead to spin precession even in the absence of an applied magnetic field (König and Martinek, 2003).

3. Andreev reflection

Andreev reflection (Andreev, 1964) is a scattering process, at an interface with a superconductor, responsible for a conversion between a dissipative quasiparticle current and a dissipationless supercurrent [see also early work by de Gennes and Saint James (1963)]. For a spin-singlet superconductor an incident electron (hole) of spin λ is reflected as a hole (electron) belonging to the opposite spin subband $\bar{\lambda}$, back to the nonsuperconducting region, while a Cooper pair is transferred to the superconductor. This is a phase-coherent scattering process in which the reflected particle carries the information about both the phase of the incident particle and the macroscopic phase of the superconductor.⁹⁶ Andreev reflection thus is responsible for a proximity effect where the phase correlations are introduced to a nonsuperconducting material (Bergeret *et al.*, 2001; Demler *et al.*, 1997; Fominov, 2003; Halterman and Valls, 2002; Izyumov *et al.*, 2002). The probability for Andreev reflection at low bias voltage ($qV \lesssim \Delta$), which is related to the square of the normal-state transmission, could be ignored for low-transparency junctions with conventional superconductors, as discussed in Sec. IV.A.1. In contrast, for high-transparency junctions (see the discussion of Sharvin con-

ductance in Sec. II.C.2), single-particle tunneling vanishes [recall Eq. (100)] at low bias and $T = 0$ and Andreev reflection is the dominant process. A convenient description is provided by the Bogoliubov-de Gennes equations (de Gennes, 1989),

$$\begin{bmatrix} H_\lambda & \Delta \\ \Delta^* & -H_{\bar{\lambda}}^* \end{bmatrix} \begin{bmatrix} u_\lambda \\ v_{\bar{\lambda}} \end{bmatrix} = E \begin{bmatrix} u_\lambda \\ v_{\bar{\lambda}} \end{bmatrix}, \quad (103)$$

and by matching the wave functions at the boundaries (interfaces) between different regions. Here H_λ is the single-particle Hamiltonian for spin $\lambda = \uparrow, \downarrow$ and $\bar{\lambda}$ denotes a spin opposite to λ (de Jong and Beenakker, 1995; Žutić and Valls, 2000). Δ is the pair potential (de Gennes, 1989), E the excitation energy and $u_\lambda, v_{\bar{\lambda}}$ are the electronlike quasiparticle and holelike quasiparticle amplitudes, respectively.⁹⁷ Griffin and Demers (1971) have solved the Bogoliubov-de Gennes equations with square or a δ -function barriers of varying strength at an N/S interface. They obtained a result that interpolates between the clean and the tunneling limits. Blonder *et al.* (1982) used a similar approach, known as the Blonder-Tinkham-Klapwijk method, in which the two limits correspond to $Z \rightarrow 0$ and $Z \rightarrow \infty$, respectively, and Z is the strength of the δ -function barrier. The transparency of this approach⁹⁸ makes it suitable for the study of ballistic spin-polarized transport and spin injection even in the absence of a superconducting region (Heersche *et al.*, 2001; Hu and Matsuyama, 2001; Hu *et al.*, 2001a; Matsuyama *et al.*, 2002).

It is instructive to note a similarity between the two-component transport in N/S junctions (for electronlike and holelike quasiparticles) and F/N junctions (for spin \uparrow, \downarrow), which both lead to current conversion, accompanied by the additional boundary resistance (Blonder *et al.*, 1982; van Son *et al.*, 1987). In the N/S junction Andreev reflection is responsible for the conversion between the normal and the supercurrent, characterized by the superconducting coherence length, while in the F/N case a conversion between spin-polarized and unpolarized current is characterized by the spin diffusion length.

For spin-polarized carriers, with different populations in two spin subbands, only a fraction of the incident electrons from a majority subband will have a minority subband partner in order to be Andreev reflected. This can be simply quantified at zero bias and $Z = 0$, in terms of the total number of scattering channels (for each $k_{||}$)

⁹⁷ Equation (103) can be simply modified to include the spin flip and spin-dependent interfacial scattering (Žutić and Das Sarma, 1999).

⁹⁸ A good agreement (Yan *et al.*, 2000) was obtained with the more rigorous nonequilibrium Keldysh technique (Keldysh, 1964; Rammer and Smith, 1986), for an illustration of how such a technique can be used to study spin-polarized transport in a wide range of heterojunctions see Mélin and Feinberg (2002); Zeng *et al.* (2003).

junctions (Pederson and Vernon, Jr., 1967), CPP multilayers (Lenczowski *et al.*, 1994), and conventional MTJ's (Moodera *et al.*, 1996; Rzchowski and Wu, 2000).

⁹⁶ For instructive reviews see Lambert and Raimondi (1998); Pannetier and Courtois (2000).

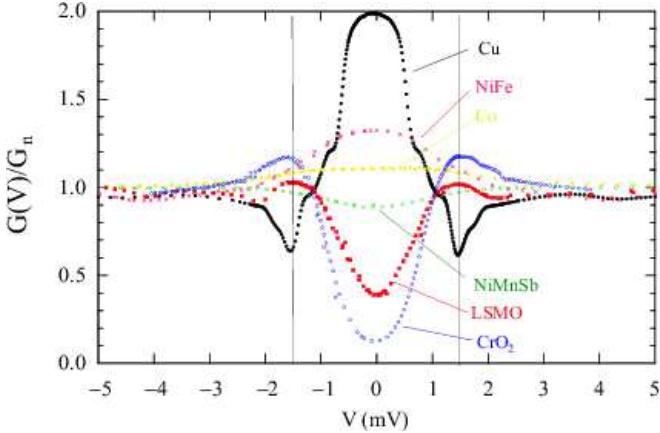


FIG. 24 The differential conductance for several spin-polarized materials, showing the suppression of Andreev reflection with increasing P_G . The vertical lines denote the bulk superconducting gap for Nb: $\Delta(T = 0) = 1.5$ meV. Note that NiMnSb, one of the Heusler alloys originally proposed as half-metallic ferromagnets (de Groot *et al.*, 1983b), shows only partial spin polarization. From Soulen Jr. *et al.*, 1998.

$N_\lambda = k_{F\lambda}^2 A / 4\pi$ at the Fermi level. Here A is the point-contact area, and $k_{F\lambda}$ is the spin-resolved Fermi wave vector. A spherical Fermi surface in the F and S regions, with no (spin-averaged) Fermi velocity mismatch, is assumed. When S is in the normal state, the zero-temperature Sharvin conductance is

$$G_{FN} = \frac{e^2}{h} (N_\uparrow + N_\downarrow), \quad (104)$$

equivalent to R_{Sharvin}^{-1} , from Eq. (47). In the superconducting state all of the N_\downarrow and only $(N_\downarrow/N_\uparrow)N_\uparrow$ scattering channels contribute to Andreev reflection across the F/S interface and transfer charge $2e$, yielding (de Jong and Beenakker, 1995)

$$G_{FS} = \frac{e^2}{h} \left(2N_\downarrow + \frac{2N_\downarrow}{N_\uparrow} N_\uparrow \right) = 4 \frac{e^2}{h} N_\downarrow. \quad (105)$$

The suppression of the normalized zero-bias conductance at $V = 0$ and $Z = 0$ (de Jong and Beenakker, 1995),

$$G_{FS}/G_{FN} = 2(1 - P_G) \quad (106)$$

with the increase in the spin polarization $P_G \rightarrow (N_\uparrow - N_\downarrow)/(N_\uparrow + N_\downarrow)$, was used as a sensitive transport technique to detect spin polarization in a point contact (Soulen Jr. *et al.*, 1998). Data are given in Fig. 24. A similar study, using a thin-film nanocontact geometry (Upadhyay *et al.*, 1998), emphasized the importance of fitting the conductance data over a wide range of applied bias, not only at $V = 0$, in order to extract the spin polarization of the F region more precisely.

The advantage of such techniques is the detection of polarization in a much wider range of materials than those which can be grown for detection in F/I/S or F/I/F

tunnel junctions. A large number of experimental results using spin-polarized Andreev reflection has since been reported (Bourgeois *et al.*, 2001; Ji *et al.*, 2001; Nadgorny *et al.*, 2001; Panguluri *et al.*, 2003b; Parker *et al.*, 2002), including the first direct measurements (Braden *et al.*, 2003; Panguluri *et al.*, 2003a, 2004) of the spin polarization in (Ga,Mn)As and (In,Mn)Sb.⁹⁹ However, for a quantitative interpretation of the measured polarization, important additional factors (similar to the limitations discussed for the application of Jullière's formula in Sec. IV.A.2) need to be incorporated in the picture provided by Eq. (106). For example, the Fermi surface may not be spherical [see the discussion of Mazin (1999), specifying what type of spin polarization is experimentally measured and also that of Xia *et al.* (2002)]. The roughness or the size of the F/S interface may lead to a diffusive component of the transport (Fal'ko *et al.*, 1999; Jedema *et al.*, 1999; Mazin *et al.*, 2001). As a caution concerning the possible difficulties in analyzing experimental data, we mention some subtleties that arise even for the simple model of a spherical Fermi surface used to describe both F and S regions. Unlike charge transport in N/S junctions (Blonder and Tinkham, 1983) in a Griffin-Demers-Blonder-Tinkham-Klapwijk approach, Fermi velocity mismatch between the F and the S regions, does not simply increase the value of effective Z . Specifically, at $Z = V = 0$ and normal incidence it is possible to have perfect transparency even when all the Fermi velocities differ, satisfying $(v_{F\uparrow} v_{F\downarrow})^{1/2} = v_S$, where v_S is the Fermi velocity in a superconductor (Žutić and Das Sarma, 1999; Žutić and Valls, 1999, 2000). In other words, unlike in Eq. (106), the spin polarization (nonvanishing exchange energy) can *increase* the subgap conductance, for fixed Fermi velocity mismatch. Conversely, at a fixed exchange energy, an increase in Fermi velocity mismatch could increase the subgap conductance.¹⁰⁰ In a typical interpretation of a measured conductance, complications can then arise in trying to disentangle the influence of parameters Z , P_G , and Fermi velocity mismatch from the nature of the point contacts (Kikuchi *et al.*, 2002) and the role of inelastic scattering (Auth *et al.*, 2003). Detection of P in HTSC's is even possible with a large barrier or a vacuum between the F and S regions, as proposed by Wang and Hu (2002) using resonant Andreev reflection and a *d*-wave superconductor.¹⁰¹

⁹⁹ Similar measurements were also suggested by Žutić and Das Sarma (1999) to yield information about the FSm/S interface. A more complete analysis should also quantify the effects of spin-orbit coupling.

¹⁰⁰ Similar results were also obtained when F and S region were separated with a quantum dot (Feng and Xiong, 2003; Zeng *et al.*, 2003; Zhu *et al.*, 2002) and even in a 1D tight-binding model with no spin polarization (Affleck *et al.*, 2000).

¹⁰¹ Interference effects between the quasi-electron and quasi-hole scattering trajectories that feel pair potentials of different sign lead to a large conductance near zero bias, even at large interfacial barrier (referred to as a zero-bias conductance peak in

Large magnetoresistive effects are predicted for crossed Andreev reflection (Deutscher and Feinberg, 2000), when the two F regions, separated within the distance of the superconducting coherence length,¹⁰² are on the same side of the S region. Such structures have also been theoretically studied to understand the implications of nonlocal correlations (Apinaryan and Mélin, 2002; Mélin and Feinberg, 2002).

4. Spin-polarized drift and diffusion

Traditional semiconductor devices such as field-effect transistors, bipolar diodes and transistors, or semiconductor solar cells rely in great part on carriers (electrons and holes) whose motion can be described as drift and diffusion, limited by carrier recombination. In inhomogeneous devices where charge buildup is rule, the recombination-limited drift-diffusion is supplied by Maxwell's equations, to be solved in a self-consistent manner. Many proposed spintronic devices as well as experimental settings for spin injection (see Sec. II) can be described by both carrier and spin drift and diffusion, limited by carrier recombination and spin relaxation (Fabian *et al.*, 2002b; Žutić *et al.*, 2002). In addition, if spin precession is important for device operation, spin dynamics need to be explicitly incorporated into the transport equations (Qi and Zhang, 2003). Drift of the spin-polarized carriers can be due not only to the electric field, but also to magnetic fields. We illustrate spin-polarized drift and diffusion on the transport model of spin-polarized bipolar transport, where bipolar refers to the presence of electrons and holes, not spin up and down. A spin-polarized unipolar transport can be obtained as a limiting case by setting the electron-hole recombination rate to zero and considering only one type of carrier (either electrons or holes).

Consider electrons and holes whose density is commonly denoted here as c (for carriers), moving in the electrostatic potential ϕ which comprises both the external bias V and the internal built-in fields due to charge inhomogeneities. Let the equilibrium spin splitting of the carrier band be $2q\zeta_c$. The spin λ resolved carrier charge-current density is (Žutić *et al.*, 2002)

$$\mathbf{j}_{c\lambda} = -q\mu_{c\lambda}c_\lambda\nabla\phi \pm qD_{c\lambda}\nabla c_\lambda - q\lambda\mu_{c\lambda}c_\lambda\nabla\zeta_c, \quad (107)$$

where μ and D stand for mobility and diffusion coefficients, the upper sign is for electrons and the lower sign is for holes. The first term on the right hand side describes drift caused by the total electric field, the second term represents diffusion, while the last term stands for magnetic drift—carrier drift in inhomogeneously split

bands.¹⁰³ More transparent are the equations for the total charge, $j = j_\uparrow + j_\downarrow$, and spin, $j_s = j_\uparrow - j_\downarrow$, current densities:

$$\mathbf{j}_c = -\sigma_c\nabla\phi - \sigma_{sc}\nabla\zeta_c \pm qD_c\nabla c \pm qD_{sc}\nabla s_c, \quad (108)$$

$$\mathbf{j}_{sc} = -\sigma_{sc}\nabla\phi - \sigma_c\nabla\zeta_c \pm qD_{sc}\nabla c \pm qD_c\nabla s_c, \quad (109)$$

where the carrier density $c = c_\uparrow + c_\downarrow$ and spin $s_c = c_\uparrow - c_\downarrow$, and we introduced the carrier charge and spin conductivities $\sigma_c = q(\mu_{cc} + \mu_{sc}s_c)$ and $\sigma_{sc} = q(\mu_{sc}c + \mu_{cs}c)$, where $\mu_c = (\mu_{c\uparrow} + \mu_{c\downarrow})/2$ and $\mu_{cs} = (\mu_{c\uparrow} - \mu_{c\downarrow})/2$ are charge and spin mobilities, and similarly for the diffusion coefficients. Equation (108) describes the spin-charge coupling in bipolar transport in inhomogeneous magnetic semiconductors. Spatial variations in spin density can cause charge currents. Similarly, it follows from Eq. (109) that spatial variations in carrier densities can lead to spin currents.

Steady-state carrier recombination and spin relaxation processes are described by the continuity equations for the spin-resolved carrier densities:

$$\nabla \cdot \frac{\mathbf{j}_{c\lambda}}{q} = \pm w_{c\lambda}(c_\lambda\bar{c} - c_{\lambda 0}\bar{c}_0) \pm \frac{c_\lambda - c_{-\lambda} - \lambda\tilde{s}_c}{2\tau_{sc}}. \quad (110)$$

Here w is the spin-dependent recombination rate, the bar denotes a complementary carrier ($\bar{n} = p$, for example), τ_{sc} is the spin relaxation time of the carrier c (not to be confused with the single spin decoherence time discussed in Sec. III.A.1), and $\tilde{s}_c = P_{c0}c$ is the nonequilibrium spin density, which appears after realizing that spin relaxation equilibrates spin while preserving carrier density. Finally, the set of equations is completed with Poisson's equation,

$$\varepsilon\Delta\phi = -\rho, \quad (111)$$

connecting the electric field and charge $\rho = q(p - n + N_d - N_a)$, where N_d and N_a are the donor and acceptor densities, respectively, and ε is the dielectric constant.

In many important cases Eqs. (107), (110), and (111) need to be solved self-consistently, which usually requires numerical techniques (Žutić *et al.*, 2002). In some cases it is possible to extract the relevant physics in limiting cases analytically, usually neglecting electric field or magnetic drift. In unipolar spin-polarized transport one does not need to consider carrier recombination. It also often suffices to study pure spin diffusion, if the built-in electric fields are small. Unipolar spin-polarized transport in inhomogeneous systems in the presence of electric fields was analyzed by Fabian *et al.* (2002b); Martin (2003); Pershin and Privman (2003a); Yu and Flatté (2002a,b). Spin-polarized drift and diffusion in model GaAs quantum wires was studied by Sogawa *et al.* (2000), while ramifications of magnetic drift for unipolar transport were

Sec. IV.A.1).

¹⁰² Recent theoretical findings suggest that the separation should not exceed the Fermi wavelength (Yamashita *et al.*, 2003b).

¹⁰³ Equation (107) can be viewed as the generalization of the Silsbee-Johnson spin-charge coupling (Heide, 2001; Johnson and Silsbee, 1987; Wegrowe, 2000) to bipolar transport and to systems with spatially inhomogeneous charge density.

studied by Fabian *et al.* (2002b); Martin (2003). Bipolar transport in the presence of electrical drift and/or diffusion has been studied by Beck *et al.* (2002); Fabian *et al.* (2002b); Flatté and Byers (2000); Zutić *et al.* (2002). Transient dynamics of spin drift and diffusion was considered by Fabian and Das Sarma (2002). Recently an interesting study (Saikin *et al.*, 2003) was reported on a Monte-Carlo simulation of quantum-mechanical spin dynamics limited by spin relaxation, in which quasiclassical orbital transport was carried out for the in-plane transport in III-V heterostructures where spin precession is due to bulk and structure inversion asymmetry (see Sec. III.B.2).

B. Materials considerations

Nominally highly spin-polarized materials, as discussed in the previous sections, could provide both effective spin injection into nonmagnetic materials and large MR effects, important for nonvolatile applications. Examples include half-metallic oxides such as CrO_2 , Fe_3O_4 , CMR materials, and double perovskites (Kobayashi *et al.*, 1998) [for reviews of half-metallic materials see (Fang *et al.*, 2002; Pickett and Moodera, 2001)]. Ferromagnetic semiconductors (Nagaev, 1983), known since CrBr_3 (Tsubokawa, 1960), have been demonstrated to be highly spin polarized. However, more recent interest in ferromagnetic semiconductors was spurred by the fabrication of (III,Mn)V compounds.¹⁰⁴ After the initial discovery of (In,Mn)As (Munekata *et al.*, 1989, 1991; Ohno *et al.*, 1992), most of the research has focused on (Ga,Mn)As (Hayashi *et al.*, 1997; Ohno *et al.*, 1996; Van Esch *et al.*, 1997). In contrast to (In,Mn)As and (Ga,Mn)As with high carrier density ($\sim 10^{20} \text{ cm}^{-3}$), a much lower carrier density in (Zn,Cr)Te (Saito *et al.*, 2002), a II-VI ferromagnetic semiconductor with Curie temperature T_C near room temperature (Saito *et al.*, 2003), suggests that transport properties can be effectively controlled by carrier doping. Most of the currently studied ferromagnetic semiconductors are p-doped with holes as spin-polarized carriers, which typically leads to lower mobilities and shorter spin relaxation times than in n-doped materials. It is possible to use selective doping to substantially increase T_C , as compared to the uniformly doped bulk ferromagnetic semiconductors (Nazmul *et al.*, 2003).

Early work on (Ga,Mn)As (De Boeck *et al.*, 1996) showed the low solubility of Mn and the formation of magnetic nanoclusters characteristic of many subsequent compounds and different magnetic impurities. The presence of such nanoclusters often complicates accurate determination of T_C as well as of whether the compound

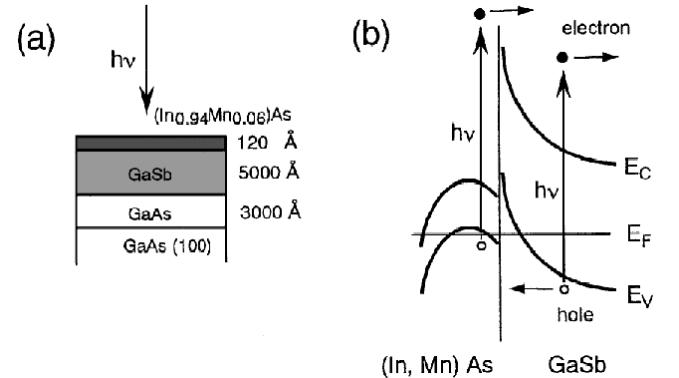


FIG. 25 Photoinduced ferromagnetism in a (In,Mn)As/GaSb heterostructure: (a) light-irradiated sample displaying photoinduced ferromagnetism—direction of light irradiation is shown by an arrow; (b) band-edge profile of (In,Mn)As/GaSb heterostructure. E_c , conduction band, E_v , valence band; E_F , Fermi level, respectively. From Koshihara *et al.*, 1997.

is actually in a single phase. Consequently, the reported room-temperature ferromagnetism in an increasing number of compounds reviewed by Pearton *et al.* (2003) is not universally accepted. Conclusive evidence for intrinsic ferromagnetism in semiconductors is highly nontrivial. For example, early work reporting ferromagnetism even at nearly 900 K in La-doped CaBa_6 (Ott *et al.*, 2000; Tromp *et al.*, 2001; Young *et al.*, 1999), was later revisited suggesting extrinsic effect (Bennett *et al.*, 2003). It remains to be understood what the limitations are for using extrinsic ferromagnets and, for example, whether they can be effective spin injectors.

A high T_C and almost complete spin polarization in bulk samples are alone not sufficient for successful applications. Spintronic devices typically rely on inhomogeneous doping, structures of reduced dimensionality, and/or structures containing different materials. Interfacial properties, as discussed in the previous sections, can significantly influence the magnitude of magnetoresistive effects¹⁰⁵ and the efficiency of spin injection. Doping properties and possibility of fabricating a wide range of structures allow spintronic applications beyond MR effects, for example, spin transistors, spin lasers, and spin-based quantum computers (Sec. IV.F). Materials properties of hybrid F/Sm heterostructures, relevant to device applications, were reviewed by Samarth *et al.* (2003).

Experiments in which the ferromagnetism is induced optically (Koshihara *et al.*, 1997; Oiwa *et al.*, 2002; Wang *et al.*, 2003b) and electrically (Ohno *et al.*, 2000a; Park *et al.*, 2002) provide a method for distinguishing the carrier-induced ferromagnetism, based on the exchange

¹⁰⁴ Ferromagnetic order with Mn-doping was obtained previously, for example, in (Sn,Mn)Te (Escorne *et al.*, 1974), (Ge,Mn)Te (Cochrane *et al.*, 1974) and (Pb,Sn,Mn)Te (Story *et al.*, 1986).

¹⁰⁵ In magnetic multilayers GMR is typically dominated by interfacial scattering (Parkin, 1993), while in MTJ's it is the surface rather than the bulk electronic structure which influences the relevant spin polarization.

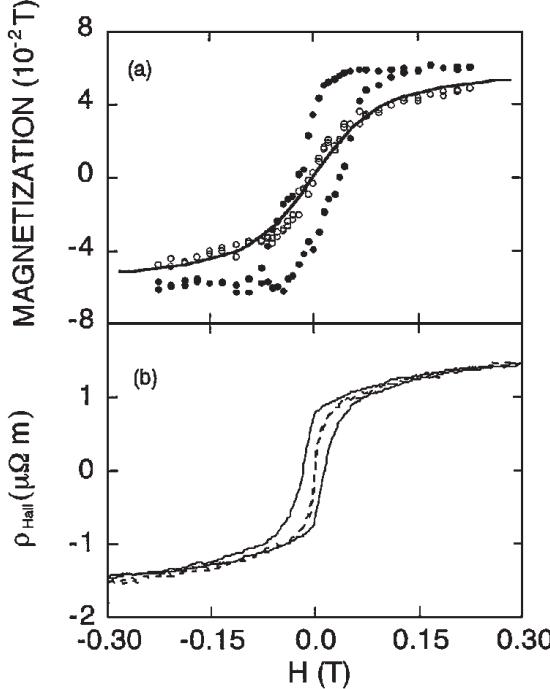


FIG. 26 Magnetization curves for $(\text{In},\text{Mn})\text{As}/\text{GaSb}$ at 5 K observed before (open circles) and after (solid circles) light irradiation. Solid line show a theoretical curve. (b) Hall resistivity at 5 K before (dashed lines) and after (solid lines) light irradiation. From Koshihara *et al.*, 1997.

interaction between the carrier and the magnetic impurity spins, from ferromagnetism that originates from magnetic nanoclusters. Such experiments also suggest a possible nonvolatile multifunctional devices with tunable, optical, electrical, and magnetic properties. Comprehensive surveys of magneto-optical materials and applications, not limited to semiconductors, are given by Sugamo and Kojima (Eds.) (2000); Zvezdin and Kotov (1997).

Photoinduced ferromagnetism was demonstrated by Koshihara *et al.* (1997) in p-(In,Mn)As/GaSb heterostructure, shown in Figs. 25, and 26. Unpolarized light penetrates through a thin (In,Mn)As layer and is absorbed in the GaSb layer. A large band bending across the heterostructures separates, by a built-in field, electrons and holes. The excess holes generated in a GaSb layer are effectively transferred to the p-doped (In,Mn)As layer where they enhance the ferromagnetic spin exchange among Mn ions, resulting in a paramagnetic-ferromagnetic transition.

The increase in magnetization, measured by a SQUID, is shown in Fig. 26(a) and in Fig. 26(b) the corresponding Hall resistivity

$$\rho_{\text{Hall}} = R_0 B + R_S M, \quad (112)$$

is shown, where the R_0 is the ordinary and R_S the anomalous Hall coefficient, respectively. Typical for $(\text{III},\text{Mn})\text{V}$

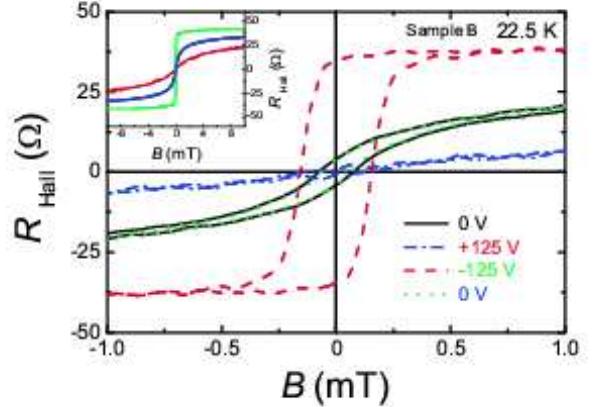


FIG. 27 Electric-field control of ferromagnetism. R_{Hall} vs field curves under three different gate biases. Application of $V_G=0$, $+125$, and -125 V results in a qualitatively different field dependence of R_{Hall} measured at 22.5 K (sample B): almost horizontal dash-dotted line, paramagnetic response are partially depleted from the channel ($V_G=+125$ V); dashed lines, clear hysteresis at low fields (<0.7 mT) as holes are accumulated in the channel ($V_G=-125$ V); solid line, R_{Hall} curve measured at $V_G=0$ V before application of ± 125 V, dotted line, R_{Hall} after application of ± 125 V. Inset, the same curves shown at higher magnetic fields. From Ohno *et al.*, 2000a.

compounds, ρ_{Hall} is dominated by the anomalous contribution, $\rho_{\text{Hall}} \propto M$.

A different type of photoinduced magnetization was measured in ferromagnetic $(\text{Ga},\text{Mn})\text{As}$.¹⁰⁶ In a Faraday geometry (recall II.D.3), by changing the polarization of a circularly polarized light, one can modulate the Hall resistance and thus the induced magnetization by up to 15% of the saturation value (Oiwa *et al.*, 2002). Additional experiments on photoinduced magnetization rotation (Munekata *et al.*, 2003; Oiwa *et al.*, 2003) suggest that the main contribution of carrier spin to such rotation is realized by generating an effective magnetic field through the p - d exchange interaction, rather than by spin-transfer torque as discussed in Secs. I.B.1 and V (Moriya *et al.*, 2003). In GaAs-Fe composite films an observation of room temperature photoenhanced magnetization was used to demonstrate that a magnetic force can be changed by light illumination (Shinshi *et al.*, 2003).

Electrically induced ferromagnetism was realized by applying gate voltage V_G to change the hole concentration in $d=5\text{nm}$ thick (In,Mn)As used as a magnetic channel in a metal-insulator semiconductor FET struc-

¹⁰⁶ Previous studies in paramagnetic $(\text{II},\text{Mn})\text{VI}$ materials have shown that nonequilibrium spin-polarized carriers can change the orientation of magnetic spins in $(\text{Hg},\text{Mn})\text{Te}$ (Krenn *et al.*, 1989, 1985) and in $(\text{Cd},\text{Mn})\text{Te}$ (Awschalom *et al.*, 1987).

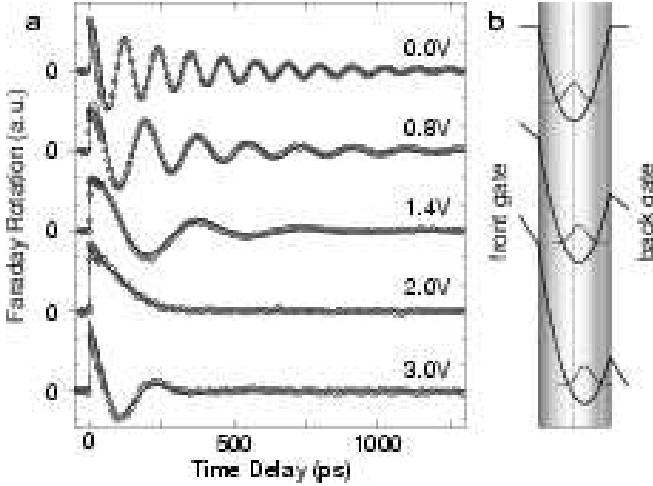


FIG. 28 Voltage-controlled spin precession: (a) time-resolved Kerr rotation measurements of electron spin precession in a quantum well at different gate voltages V_G with Al concentration of 7% at 5 K and $B=6$ T; displacement of the electron wave function towards the back gate into regions with more Al concentration as a positive voltage V_G is applied between back and front gate; leading to an increase of g . At $V_G=2$ V, no precession is observed, corresponding to $g=0$. From Salis *et al.*, 2001b.

ture. Below a metal gate and an insulator the (In,Mn)As channel was grown on top of a InAs/(Al,Ga)Sb/AlSb and GaAs substrate. In Fig. 27, the corresponding data for $R_{\text{Hall}} = \rho_{\text{Hall}}/d \propto M$ [recall Eq. (112), show that the ferromagnetism can be switched on and off, as an electric analog of the manipulation of M from Fig. 26. Subsequent work by Park *et al.* (2002) showed that in MnGe ferromagnetism can be manipulated at higher temperature and at significantly lower gate voltage (at ~ 50 K and ~ 1 V). The combination of light and electric-field control of ferromagnetism was used in modulation-doped p-type (Cd,Mn)Te QW (Boukari *et al.*, 2002). It was demonstrated that illumination by light in $p-i-n$ diodes would enhance the spontaneous magnetization, while illumination in $p-i-p$ structures would destroy ferromagnetism.

In semiconductors g factors, which determine the spin splitting of carrier bands (and consequently influence the spin dynamics and spin resonance), can be very different from the free-electron value. With strong spin-orbit coupling in narrow-band III-V's they are ≈ -50 for InSb and ≈ -15 for InAs, while, as discussed in Sec. II.D.3 the doping with magnetic impurities can give even $|g^*| \sim 500$. Manipulation of the g factor in a GaAs/AlGaAs quantum well (QW) in Fig. 28, relies on the results for a bulk $\text{Al}_x\text{Ga}_{1-x}\text{As}$, the variation of Al concentration changes the g factor (Chadi *et al.*, 1976; Weisbuch and Hermann, 1977) $g=-0.44$ for $x=0$ and $g=0.40$ for $x=0.3$. Related experiments on modulation-doped GaAs/Al_{0.3}Ga_{0.7}As have shown that by applying V_G one can shift the electron wave function in the QW and produce $\sim 1\%$ change in the

g factor (Jiang and Yablonovitch, 2001). Subsequently, in an optimized $\text{Al}_x\text{Ga}_{1-x}\text{As}$ quantum well, where x varied gradually across the structure, much larger changes were measured—when V_G is changed, the electron wave function efficiently samples different regions with different g factors (Salis *et al.*, 2001b). Figure 28(a) gives the time-resolved Kerr rotation data (the technique is discussed in Sec. III) data which can be described as $\propto \exp(-\Delta t/T_2^*) \cos(\Omega \Delta t)$, where Δt is the delay time between the circularly polarized pump and linearly polarized probe pulses, T_2^* is the transverse electron spin lifetime with inhomogeneous broadening, and the angular precession frequency $\Omega = \mu_B g B / \hbar$ can be used to determine the g factor. It is also possible to manipulate g factors dynamically using time-dependent V_G (Kato *et al.*, 2003). The anisotropy of g factor (g tensor) allows voltage control of both the magnitude and the direction of the spin precession vector Ω .

C. Spin filters

Solid state spin filtering (recall the similarity with spin injection from Sec II.C.1) was first realized in N/F/N tunneling. It was shown by Esaki *et al.* (1967) that the magnetic tunneling through (ferro)magnetic semiconductor Eu chalcogenides (Kasuya and Yanase, 1968; Nagaev, 1983; von Molnár and Methfess, 1967), such as EuSe¹⁰⁷ and EuS¹⁰⁸ could be modified by an applied magnetic field. The change in $I - V$ curves in the N/F/N structure, where N is a normal metal and F is a ferromagnet, was explained by the influence of the magnetic field on the height of the barrier formed at the N/F interface (for EuSe, the barrier height was lowered by 25% at 2 T). The large spin splitting of the Eu chalcogenides was subsequently employed in the absence of applied field with EuS (Moodera *et al.*, 1988) and nearly 100% spin polarization was reached at $B=1.2$ T with EuSe (Moodera *et al.*, 1993). These spin-filtering properties of the Eu chalcogenides, used together with one-electron quantum dots, were proposed as the basis for a method to convert single spin into single charge measurements¹⁰⁹ and provide an important ingredient in realizing a quantum computer (DiVincenzo, 1999), see Sec. IV.F.

Zeeman splitting in semiconductor heterostructures and superlattices (enhanced by large g factors) (Egues, 1998; Guo *et al.*, 2001), in quantum dots (Borda *et al.*, 2003; Deshmukh and Ralph, 2002; Recher *et al.*, 2000), and nanocrystals (Efros *et al.*, 2001) provide effective spin filtering and spin-polarized currents. Predicted

¹⁰⁷ At zero magnetic field EuSe is an antiferromagnet, and at moderate fields it becomes a ferromagnet with $T_C \approx 5$ K.

¹⁰⁸ At zero magnetic field, exchange splitting of a conduction band in bulk EuS is ≈ 0.36 eV (Hao *et al.*, 1990).

¹⁰⁹ This method could already be realized using single-electron transistors or quantum point contacts.

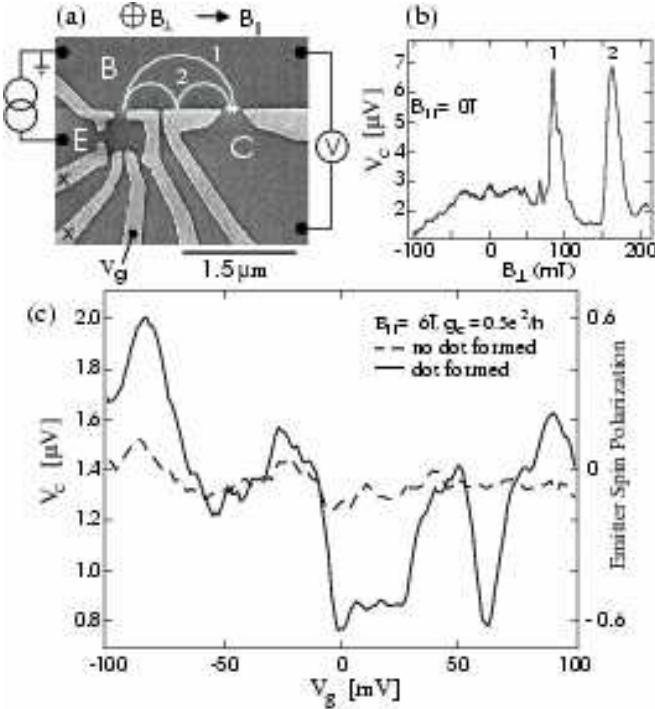


FIG. 29 Mesoscopic spin filter: (a) micrograph and circuit showing the polarizer-analyzer configuration used in the experiment of Folk *et al.* (2003). The emitter (E) can be formed into either a quantum dot or a quantum point contact (QPC). The collector (C), is a single point contact. Electrons are focused from E to C through the base region (B), using a small perpendicular magnetic field. Gates marked with “x” are left undepleted when E is operated as a QPC; (b) base-collector voltage (V_C) showing two focusing peaks; (c) focusing peak height at $B_{\parallel}=6$ T with spin-selective collector QPC conductance ($g_C=0.5e^2/h$), comparing E as QPC at $2e^2/h$ (dashed curve) and E as a quantum dot with both leads at $2e^2/h$ (solid curve). Adapted from Folk *et al.*, 2003.

quantum size effects and resonance tunneling (Duke, 1969) also have their spin-dependent counterparts. The structures studied are typically double-barrier resonant tunneling diodes (for an early spin-unpolarized study see Tsu and Esaki (1973)), with either Zeeman splitting or using ferromagnetic materials, in which spin filtering can be tuned by an applied bias.¹¹⁰

Several other realizations of spin filtering have been investigated, relying on spin-orbit coupling,¹¹¹ or hot-electron transport across ferromagnetic regions,¹¹² dis-

cussed in more detail in Sec. IV.E.3. A choice of particular atomically ordered F/Sm interfaces was suggested to give a strong spin-filtering effect (Kirczenow, 2001), limited by the spin-orbit coupling and interfacial spin-flip scattering.

Mesoscopic spin filters have also been suggested (Avdonin *et al.*, 2003; Frustaglia *et al.*, 2001; Ionicioiu and D’Amico, 2003; Joshi *et al.*, 2001) and we discuss here a particular realization. In an applied magnetic field two quantum point contacts (QPC), an emitter (E) and a collector (C), fabricated on top of a high-mobility 2DEG in GaAs/AlGaAs, can act as spin polarizer and analyzer (Potok *et al.*, 2002).¹¹³ In a ballistic regime and at $T = 70$ mK (mean free path ≈ 45 μm \gg sample size ≈ 1.5 μm) magnetic focusing¹¹⁴ with B_{\perp} results in base-collector voltage peaks, when the separation of the two QPCs is an even multiple of the cyclotron radius m^*v_F/eB_{\perp} , where m^* is the effective mass, and v_F the Fermi velocity. These results are illustrated in Figs. 29(a) and (b) on a slightly modified structure (Folk *et al.*, 2003) where, by applying a gate voltage, one can cause the emitter to form either a quantum dot or QPC. Effective spin filtering, due to the large in-plane field B_{\parallel} , can be tuned by the gate voltage which changes the conductance of QPC. The resulting effect of spin filtering modifies the collector voltage V_C (Potok *et al.*, 2002),

$$V_C = \alpha(h/2e^2)I_E(1 + P_{I_E}P_{T_C}), \quad (113)$$

where $0 < \alpha < 1$ parameterizes the imperfections in focusing, P_{I_E} and P_{T_C} are the spin polarization [recall Eq. (3)] of the emitter current I_E , and the collector transmission coefficient T_C is related to the collector conductance by $g_C = (2e^2/h)T_C$. In Eq. (113) we note a recurring form for a spin-valve effect. The measured signal involves the product of two different spin polarizations, for example, similar to TMR in Eq. (2) or to spin-charge coupling due to nonequilibrium spin [recall Eqs. (43) and (114)]. Another mesoscopic spin filter with few-electron quantum dot (GaAs/AlGaAs-based) was used to demonstrate a nearly complete spin polarization which could be reversed by adjusting gate voltages (Hanson *et al.*, 2003a).

D. Spin diodes

Spin diodes are inhomogeneous two-terminal devices whose electronic or optical properties depend on the spin

¹¹⁰ See, for example, (Aleiner and Lyanda-Geller, 1991; Brehmer *et al.*, 1995; Giazotto *et al.*, 2003; Mendez *et al.*, 1998; Ohno, 1998; Petukhov, 1998; Petukhov *et al.*, 2002; Slobodskyy *et al.*, 2003; Ting and Cartoixa, 2002; Vurgaftman and Meyer, 2003).

¹¹¹ These include the work of (de Andrada e Silva and La Rocca, 1999; Governale *et al.*, 2002; Kiselev and Kim, 2001; Koga *et al.*, 2002b; Perel’ *et al.*, 2003; Voskoboynikov *et al.*, 1998, 1999).

¹¹² See (Cacho *et al.*, 2002; Filipe *et al.*, 1998; Monsma *et al.*, 1995;

Oberli *et al.*, 1998; Rippard and Buhrman, 2000; Upadhyay *et al.*, 1999; van Dijken *et al.*, 2002b).

¹¹³ QPC was also used to locally create and probe nonequilibrium nuclear spin in GaAs/AlGaAs heterostructures in the quantum Hall regime (Wald *et al.*, 1994).

¹¹⁴ Suggested by Sharvin (1965) as a technique to study Fermi surfaces; see also van Houten *et al.* (1989).

polarization of the carriers. Such devices were envisaged long before the emergence of spintronics. Solomon (1976), for example, proposed and demonstrated a silicon *p-n* junction whose current was modified by changing the spin polarization of the recombination centers. In a magnetic field both the mobile carriers and the recombination centers have an equilibrium spin polarization due to the Zeeman splitting. The current in a *p-n* junction depends on the recombination rate, which, in turn, depends on the relative orientation of the spin of the carriers and the centers (Lepine, 1972). The trick to modifying the current is to decrease (even saturate) the spin polarization of either the electrons or the centers by electron spin resonance. Indeed, Solomon (1976) found a variation of $\approx 0.01\%$ of the saturation current at small biases where recombination in the space-charge region dominates. Similar experiments could be used to detect nonequilibrium spin due to (potential) spin injection in Si, where optical methods are ineffective, but also in other semiconductors where electrical detection would be desirable.¹¹⁵

Several spin diodes have recently been proposed or demonstrated with the goal of either maximizing the sensitivity of the $I - V$ characteristics to spin and magnetic field, or to facilitating spin injection and its detection through semiconductor interfaces comprising a magnetic semiconductor as the injector. Magnetic tunneling diodes have been used for spin injection from a ferromagnetic to a nonmagnetic semiconductor, in p-GaMnAs/n-GaAs *p-n* junctions (Johnston-Halperin *et al.*, 2002; Kohda *et al.*, 2001; Van Dorpe *et al.*, 2003a). As discussed in Sec. II.D.3, *p-n* heterostructures have combined Cr- or Eu-based ferromagnetic semiconductors and InSb (Osipov *et al.*, 1998; Viglin *et al.*, 1997). Spin light-emitting diodes (recall Figs. 12 and 13) were employed for injecting and detecting spins in semiconductors, while resonant tunneling diodes have been demonstrated as effective spin injectors (Sec. II.D.3) and spin filters (Sec. IV.C). A magnetic unipolar diode has been proposed by Flatté and Vignale (2001) to simulate the working of ordinary diodes, but with homogeneous monopolar doping (either donors or acceptors, not both). The role of inhomogeneous doping in the *p-n* junction is played by the inhomogeneous spin splitting of the carrier band, with the spin up and spin down carriers playing roles similar to those of the electrons and holes in bipolar diodes. Si-based *p-i-n* diode sandwiched between two ferromagnetic metals was suggested to allow controlling the device perfor-

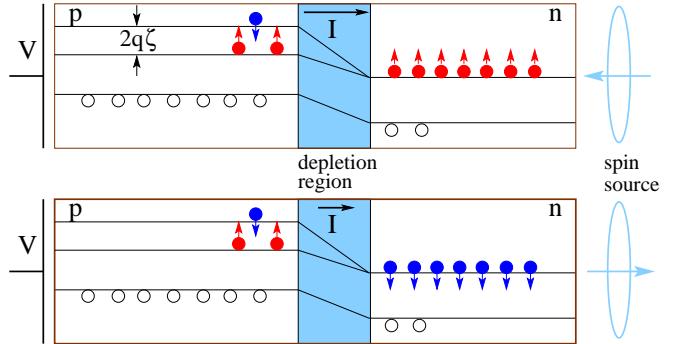


FIG. 30 Scheme of a magnetic bipolar diode. The *p*-region (left) is magnetic, indicated by the spin splitting $2q\zeta$ of the conduction band. The *n*-region (right) is nonmagnetic, but spin polarized by a spin source: Filled circles, spin-polarized electrons; empty circles, unpolarized holes. If the nonequilibrium spin in the *n*-region is oriented parallel (top figure) to the equilibrium spin in the *p*-region, large forward current flows. If the relative orientation is antiparallel (bottom), the current drops significantly. Adapted from Žutić *et al.*, 2002.

mance by an externally applied magnetic field (Dugaev *et al.*, 2003). Finally, Žutić *et al.* (2002) have proposed the magnetic bipolar diode described below.

The magnetic bipolar diode¹¹⁶ (MBD) is a *p-n* junction diode with one or both regions magnetic (Fabian *et al.*, 2002b; Žutić *et al.*, 2002). The MBD is the prototypical device of bipolar spintronics, a subfield of spintronics in which both electrons and holes take part in carrier transport, while either electrons or holes (or both) are spin polarized (see Sec. IV.A.4). Examples of nonmagnetic bipolar spintronic devices are the spin-polarized *p-n* junction (Žutić *et al.*, 2001b) and the spin solar cell (Žutić *et al.*, 2001a). These devices offer opportunities for effective spin injection, spin amplification (see Sec. II.C.3), or spin capacity—the effect of changing, by voltage, nonequilibrium spin density (Žutić *et al.*, 2001b). The advantages of magnetic bipolar spintronic devices (Fabian *et al.*, 2002a,b; Žutić *et al.*, 2002, 2003) lie in the combination of equilibrium magnetism and nonequilibrium spin and effective methods to manipulate a minority carrier population. The most useful effects of the spin-charge coupling in MBD's are the spin-voltaic and the giant-magnetoresistive effects, which are enhanced over those of metallic systems by the exponential dependence of the current on bias voltage.

A scheme of an MBD is shown in Fig. 30 (also see Fig. 8). The *p*-region is magnetic, by which we mean that it has a spin-split conduction band with the spin splitting (Zeeman or exchange) $2q\zeta \sim k_B T$. Zeeman

¹¹⁵ Spin diodes can also probe fundamental properties of electronic systems. The diode demonstrated by Kane *et al.* (1992) is based on a junction between two coplanar AlGaAs/GaAs 2DEG's, one with $\nu < 1$ and the other with $\nu > 1$, where ν is the Landau-level filling; that is, the two regions have opposite spins at the Fermi level. The current crossing such a junction, which has a diode property due to the existence of a built-in field in the contact, is accompanied by a spin flip. Interestingly, the current is also time dependent, due to the current-induced dynamic nuclear polarization.

¹¹⁶ Not to be confused with the usual magnetic diodes which are ordinary diodes in a magnetic field. The $I - V$ characteristics of such diodes, depend on the magnetic field through small orbital effects on diffusion coefficient, not through the spin effects described here.

splitting can be significantly enhanced by the large g^* factors of magnetically doped (Sec. II.C.3) or narrow-band-gap semiconductors (Sec. IV.B). Using an MBD with a ferromagnetic semiconductor slightly above its T_C is also expected to give large g^* factors. The n -region is nonmagnetic, but electrons can be spin-polarized by a spin source (circularly polarized light or magnetic electrode). The interplay between the equilibrium spin of polarization $P_{n0} = \tanh(q\zeta/k_B T)$ in the p -region, and the nonequilibrium spin source of polarization δP_n in the n -region, at the edge of the depletion layer, determines the $I - V$ characteristics of the diodes. It is straightforward to generalize these considerations to include the spin-polarized holes (Fabian *et al.*, 2002b).

The dependence of the electric current j on $q\zeta$ and δP_n was obtained by both numerical and analytical methods. Numerical calculations (Žutić *et al.*, 2002) were performed by self-consistently solving for the drift-diffusion, continuity, as well as carrier recombination and spin-relaxation equations, discussed in Sec. IV.A.4. While the numerical calculations are indispensable in the high-injection limit,¹¹⁷ valuable insight and analytical formulas can be obtained in the low-injection limit, where the Shockley theory (Shockley, 1950) for ordinary p - n junctions was generalized by Fabian *et al.* (2002b) for the magnetic case.

To illustrate the $I - V$ characteristics of MBD's, consider the low-injection limit in the configuration of Fig. 30. The electron contribution to the total electric current is (Fabian *et al.*, 2002b)

$$j_n \sim n_0(\zeta) \left[e^{qV/k_B T} (1 + \delta P_n P_{n0}) - 1 \right], \quad (114)$$

where V is the applied bias (positive for forward bias) and $n_0(\zeta) = (n_i^2/N_a) \cosh(q\zeta/k_B T)$ is the equilibrium number of electrons in the p -region, dependent on the splitting, the intrinsic carrier density n_i , and the acceptor doping N_a . Equation (114) generalizes the Silsbee-Johnson spin-charge coupling (Johnson and Silsbee, 1985; Silsbee, 1980), originally proposed for ferromagnet/paramagnet metal interfaces, to the case of magnetic p - n junctions. The advantage of the spin-charge coupling in p - n junctions, as opposed to metals or degenerate systems, is the nonlinear voltage dependence of the nonequilibrium carrier and spin densities (Fabian *et al.*, 2002b), allowing for the exponential enhancement of the

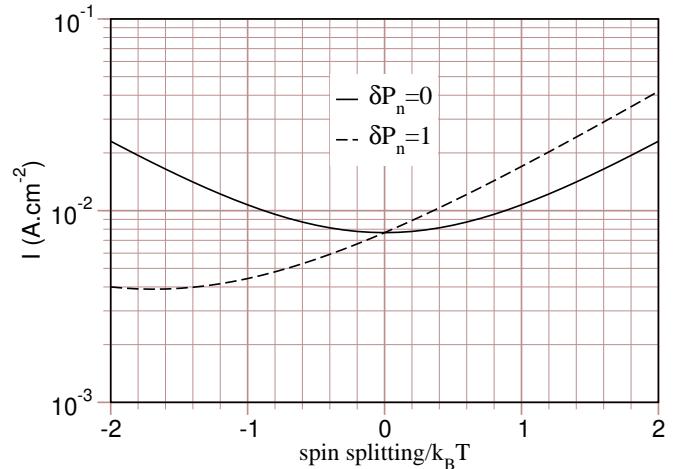


FIG. 31 Giant magnetoresistance (GMR) effect in magnetic diodes. Current/spin-splitting characteristics ($I - \zeta$) are calculated self-consistently at $V=0.8$ V for the diode from Fig. 30. Spin splitting $2q\zeta$ on the p-side is normalized to $k_B T$. The solid curve corresponds to a switched-off spin source. The current is symmetric in ζ . With spin source on (the extreme case of 100% spin polarization injected into the n -region is shown), the current is a strongly asymmetric function of ζ , displaying large GMR, shown by the dashed curve. Materials parameters of GaAs were applied. Adapted from Žutić *et al.*, 2002.

effect with increasing V . Equation (114) can be understood qualitatively from Fig. 30 (Fabian *et al.*, 2002b). In equilibrium, $\delta P_n = 0$ and $V = 0$, no current flows through the depletion layer, as the electron currents from both sides of the junction balance out. The balance is disturbed either by applying bias or by selectively populating different spin states, making the flow of one spin species greater than that of the other. In the latter case, the effective barriers for crossing of electrons from the n to the p side is different for spin up and down electrons (see Fig. 30). Current can flow even at $V = 0$ when $\delta P_n \neq 0$. This is an example of the spin-voltaic effect (a spin analog of the photo-voltaic effect), in which nonequilibrium spin causes an EMF (Žutić and Fabian, 2003; Žutić *et al.*, 2002). In addition, the direction of the zero-bias current is controlled by the relative sign of P_{n0} and δP_n .

MBD's can display an interesting GMR-like effect, which follows from Eq. (114) (Žutić *et al.*, 2002). The current depends strongly on the relative orientation of the nonequilibrium spin and the equilibrium magnetization. Figure 31 plots j , which also includes the contribution from holes, as a function of $2q\zeta/k_B T$ for both the unpolarized, $\delta P_n = 0$, and fully polarized, $\delta P_n = 1$, n -region. In the first case j is a symmetric function of ζ , increasing exponentially with increasing ζ due to the increase in the equilibrium minority carrier density $n_0(\zeta)$. In unipolar systems, where transport is due to the majority carriers, such a modulation of the current is not likely, as the majority carrier density is fixed by the density of

¹¹⁷ The small bias or low-injection limit is the regime of applied bias in which the density of the carriers injected through the depletion layer (the minority carriers) is much smaller than the equilibrium density of the majority carriers. Here and in Sec. IV.E.2 the terms majority and minority refer to the relative carrier (electron or hole) population and not to spin. The large bias or high-injection limit is the regime where the injected carrier density becomes comparable to the equilibrium density. This occurs at forward biases comparable to the built-in potential, typically 1 V.

dopants.

If $\delta P_n \neq 0$, the current will depend on the sign of $P_{n0} \cdot \delta P_n$. For parallel nonequilibrium (in the *n*-region) and equilibrium spins (in the *p*-region), most electrons cross the depletion layer through the lower barrier (see Fig. 30), increasing the current. In the opposite case of antiparallel relative orientation, electrons experience a larger barrier and the current is inhibited. This is demonstrated in Fig. 31 by the strong asymmetry in j . The corresponding GMR ratio, the difference between j for parallel and antiparallel orientations, can also be calculated analytically from Eq. (114) as $2|\delta P_n P_{n0}|/(1 - |\delta P_n P_{n0}|)$ (Fabian *et al.*, 2002b). If, for example, $|P_{n0}| = |\delta P_n| = 0.5$, the relative change is 66%. The GMR effect should be useful for measuring the spin relaxation rate of bulk semiconductors (Žutić *et al.*, 2003), as well as for detecting nonequilibrium spin in the nonmagnetic region of the *p-n* junction.¹¹⁸

Although practical MBD's are still to be fabricated and the predicted effects tested, magnetic *p-n* junctions have already been demonstrated. Indeed, Wen *et al.* (1968)¹¹⁹ were perhaps the first to show that a ferromagnetic *p-n* junction, based on the ferromagnetic semiconductor CdCr₂Se₄ doped with Ag acceptors and In donors, could act as a diode. Heavily doped p-GaMnAs/n-GaAs junctions were fabricated (Arata *et al.*, 2001; Johnston-Halperin *et al.*, 2002; Kohda *et al.*, 2001; Ohno *et al.*, 2000b; Van Dorpe *et al.*, 2003a) to demonstrate tunneling interband spin injection. Incorporation of (Ga,Mn)As layer in the intrinsic region of *p-i-n* GaAs diode was shown to lead to an efficient photodiode, in which the Mn ions function as recombination centers (Teran *et al.*, 2003). It would be interesting to see such devices combined with a spin injector in the bulk regions. Recently, Tsui *et al.* (2003) have shown that the current in p-CoMnGe/n-Ge magnetic heterojunction diodes can indeed be controlled by magnetic field. To have functioning MBD's at room temperature, and to observe the above predicted phenomena, several important challenges have to be met:

(i) Zeeman or exchange splitting needs to be sufficiently large to provide equilibrium spin polarization, $\gtrsim 1 - 10\%$. This may be difficult at room temperature, unless the effective *g* factor is ~ 100 at $B \sim 1$ T (Sec. II.D.3). The use of ferromagnetic semiconductors is limited by their T_C (Sec. IV.B).

(ii) For a strong spin-charge coupling [recall the discussion of Eq. (114)] a nondegenerate carrier density is desirable, which, while likely in (Zn,Cr)Te, is not easily realized in many other ferromagnetic semiconductors that are typically heavily doped (Sec. IV.B).

(iii) An effective integration of magnetic and nonmag-

netic structures into single devices (Samarth *et al.*, 2003) is needed.

(iv) The samples need to be smaller than the spin diffusion lengths, requiring high carrier mobility and long spin relaxation (easier to realize for spin-polarized electrons).

(v) The effects of actual device structures, such as two- and/or three-dimensional spin flow, interface contacts, spin-dependent band offsets and band bendings, strong spin relaxation in the depletion layers, etc., will need to be understood.

By combining two magnetic *p-n* junctions in series one can obtain a magnetic bipolar transistor (Sec. IV.E.2), a three terminal device which offers spin-dependent amplification.

E. Spin transistors

We review several proposals for spin transistors that have at least one semiconductor region and that aim at integrating spin and charge transport within traditional device schemes of the field-effect and junction transistors. Three important cases are discussed in detail: the Datta-Das spin field-effect transistor, the magnetic bipolar transistor, and the hot-electron spin transistor.

Various spin transistors that contain metallic (and insulating) regions have been proposed (Bauer *et al.*, 2003; Johnson, 1993a; You and Bader, 2000; Zvezdin *et al.*, 2003). There is also a large category of the spin single-electron transistors, first realized by Ono *et al.* (1996), and later investigated in (Barnaś and Fert, 1998; Ciorga *et al.*, 2002; Korotkov and Safarov, 1999; Martinék *et al.*, 2002). Spin single-electron transistors can be viewed as an extension of magnetic tunneling (see Sec. IV.A.2) to double tunnel junctions, where the Coulomb blockade becomes important (Takahashi and Maekawa, 1998). For a review of spin single-electron transistors see (Maekawa *et al.*, 2002).

1. Spin field-effect transistors

Datta and Das (1990) proposed what became the prototypical spintronic device scheme, the Datta-Das spin field-effect transistor (SFET) (see Fig. 1). The device is based on spin injection and spin detection by a ferromagnetic source and drain, and on spin precession about the built-in structure inversion asymmetry (Bychkov-Rashba) field Ω , Eq. (88), in the asymmetric, quasi-one-dimensional channel of an ordinary field-effect transistor. The attractive feature of the Datta-Das SFET is that spin-dependent device operation is controlled not by external magnetic fields, but by gate bias, which controls the spin precession rate.

The structure of the Datta-Das SFET is shown in Fig. 1. Consider a 2DEG confined along the plane of the unit vector \mathbf{n} . The precession axis of Ω lies always in

¹¹⁸ This could be a way to detect spin injection into Si, where optical detection is ineffective.

¹¹⁹ We thank M. Field for bringing this reference to our attention.

the channel plane (see Sec. III.B.2), so the results (unlike those for bulk inversion asymmetry) are insensitive to the relative orientation of \mathbf{n} and the principal crystal axes. Equation (88) determines the evolution of the expectation value for a spin perpendicular to the plane, $s_n = \mathbf{s} \cdot \mathbf{n}$, and a spin parallel to the in-plane \mathbf{k} , $s_{\parallel} = \mathbf{s} \cdot \mathbf{k}/k$:

$$ds_n/dt = 2\alpha_{BR}ks_{\parallel}, \quad ds_{\parallel}/dt = -2\alpha_{BR}ks_n, \quad (115)$$

where α_{BR} is the structure inversion asymmetry coefficient appearing in Eq. (88). The average spin component along Ω , $s_{\perp} = \mathbf{s} \cdot (\mathbf{k} \times \mathbf{n})/k$, is constant. As a result, $s_{\parallel} = s_{0\parallel} \cos(\omega t)$, where $\omega = 2\alpha_{BR}k$ and the injected spin at the source is labeled with zero. If φ is the angle between \mathbf{k} and the source-drain axis, the electron will reach the drain at time $t' = Lm_c/(\hbar k \cos \varphi)$, with the spin s_{\parallel} precessing at the angle $\phi = 2\alpha_{BR}m_c L/\hbar$. The average spin at the drain in the direction of magnetization is $s_{\parallel}(t') \cos \varphi + s_{0\perp} \mathbf{m} \cdot (\mathbf{k} \times \mathbf{n})$, so the current is modulated by $1 - \cos^2 \varphi \sin^2(\phi/2)$, the probability of finding the spin in the direction of magnetization \mathbf{m} .

Note that ϕ does not depend on the momentum (or energy) of the carriers. As the spread φ in the momenta increases, the modulation effect decreases. The largest effect is seen for $\varphi = 0$, where the current modulation factor is $\cos^2(\phi/2)$. It was therefore proposed (Datta and Das, 1990) that φ be limited by further confining the electron motion along $\varphi = 0$ using a one-dimensional channel as a waveguide. Spin modulation of the current becomes ineffective if transport is diffusive. Taking typical values (Koga *et al.*, 2002a; Nitta *et al.*, 1997) for $\hbar\alpha_{BR} \approx 1 \times 10^{-11}$ eV·m, and $m_c = 0.1m_e$, current modulation should be observable at source-drain separations of $L \gtrsim 1 \mu\text{m}$, setting the scale for ballistic transport. The device will work best with narrow-gap materials (Lommer *et al.*, 1988) like InAs, in which the structure inversion asymmetry dominates the spin precession (Das *et al.*, 1989; Luo *et al.*, 1988, 1990). Another option is using Si heterostructures, in which bulk inversion asymmetry is absent. However, the small magnitude of the spin-orbit interaction makes α_{BR} in Si probably rather weak.

The Datta-Das SFET is yet to be realized. There are at least four important difficulties in observing the proposed effects.

(i) The effective spin injection of spin-polarized carriers from the ferromagnetic source into a 2DEG is nontrivial (see Sec. II.D.4).

(ii) Ballistic spin-polarized transport should be realized through the channel with *uniform* α_{BR} by eliminating undesirable electric fields due to interface inhomogeneities.

(iii) The parameter α_{BR} should be effectively controllable by the gate.

(iv) The structure inversion asymmetry should dominate over the bulk inversion asymmetry, and the spin precession rate must be large enough ($\hbar\alpha_{BR} \gtrsim 10^{-11}$ eV·m) to allow at least a half precession during the ballistic transport.

These four factors present a great challenge to fabricating a Datta-Das SFET at room temperature, limiting the design to special materials and very clean interfaces. However, the modulation of α_{BR} by biasing voltage (iii) has been already convincingly demonstrated in $\text{In}_{0.53}\text{Ga}_{0.47}\text{As}/\text{In}_{0.52}\text{Al}_{0.48}\text{As}$ QW's (Grundler, 2000; Hu *et al.*, 1999; Nitta *et al.*, 1997) [for GaAs/AlGaAs 2DEG see also (Miller *et al.*, 2003)]. Initial experimental investigations of magnetoresistance in the Datta-Das SFET systems were performed by Gardelis *et al.* (1999). Recently spin precession in the Datta-Das SFET, including the bulk inversion asymmetry, was investigated by (Winkler, 2004) using $k \cdot p$ model calculations. It is not surprising that the conductance through the transistor, in the present orientation-dependent bulk inversion asymmetry, depends rather strongly on the crystallographic orientation of the two-dimensional channel (Lusakowski *et al.*, 2003). For more discussion of the Dresselhaus bulk inversion asymmetry and the Bychkov-Rashba structure asymmetry see Sec. III.B.2.b.

The Datta-Das SFET has generated great interest in mesoscopic spin-polarized transport in the presence of structure inversion asymmetry. Model calculations using the tight-binding formulation of H_{SIA} (recall Sec. II.B.2) were reported by Pareek and Bruno (2002). Further theoretical investigations on the theme of the Datta-Das spin transistor can be found in Matsuyama *et al.* (2002); Nikolić and Freericks (2001) and in an extensive review by Bournel (2000). Distinct SFET's have also been suggested, even in the absence of ferromagnetic regions which are replaced by a rotating external magnetic field of uniform strength (Wang *et al.*, 2003a). Ciuti *et al.* (2002b) proposed a ferromagnetic-oxide-semiconductor transistor, with a nonmagnetic source and drain, but with two ferromagnetic gates in series above the base channel. The relative orientation of the gates' magnetization leads to magnetoresistance effects. An SFET that can operate in the diffusive regime, in the presence of both bulk and structure inversion asymmetry, has been considered by Schliemann *et al.* (2003).

2. Magnetic bipolar transistor

The magnetic bipolar transistor (MBT) is a bipolar transistor with spin-split carrier bands and, in general, an injected spin (Fabian *et al.*, 2002a; Fabian and Žutić, 2004; Fabian *et al.*, 2004). A related device structure was already proposed by Gregg *et al.* (1997) in a push for silicon-based spintronics. In this proposal (also called SPICE for spin-polarized injection current emitter) the semiconductors have no equilibrium spin, while the spin source is provided by a ferromagnetic spin injector attached to the emitter, and another ferromagnetic metal, a spin detector, is attached to the base/collector junction to modulate the current flow. In both configurations the aim is to control current amplification by spin and magnetic field.

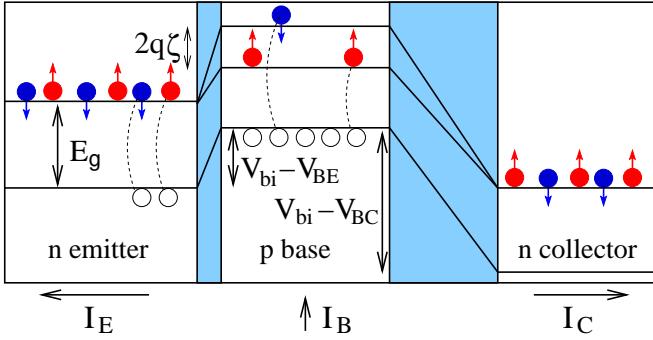


FIG. 32 Scheme of an n - p - n magnetic bipolar transistor with magnetic base (B), nonmagnetic emitter (E), and collector (C). Conduction and valence bands are separated by the energy gap E_g . The conduction band has a spin splitting $2q\zeta$, leading to equilibrium spin polarization $P_{B0} = \tanh(q\zeta/k_B T)$. Carriers and depletion regions are represented as in Fig. 30. In the so called forward active regime, where the transistor can amplify currents, the E-B junction is forward biased (here with voltage $V_{BE} > 0$ lowering the built-in potential V_{bi}), while the B-E junction is reverse biased ($V_{BC} < 0$). The directions of the current flows are indicated. Electrons flow from E to B, where they either recombine with holes (dashed lines) or continue to be swept by the electric field in the B-E depletion layer towards C. Holes form mostly the base current, I_B , flowing to the emitter. The current amplification $\beta = I_C/I_B$ can be controlled by P_{B0} as well as by the nonequilibrium spin in E. Adapted from Fabian *et al.*, 2004.

A scheme of a particular MBT is shown in Fig. 32. Such a three-terminal device can be thought of as consisting of two magnetic p - n junctions connected in series. Materials considerations discussed in Sec. IV.D also apply to an MBT in order to provide a sufficient equilibrium polarization in a magnetic base P_{B0} . While nonmagnetic, the emitter has a nonequilibrium polarization δP_E from a spin source, similar to the magnetic diode case in Fig. 30. Only the spin polarization of electrons is assumed. Applying the generalized Shockley theory to include spin effects (Fabian *et al.*, 2002b), a theory of MBT was developed by Fabian *et al.* (2002a); Fabian and Žutić (2004). Later, simplified schemes of MBT [not including the effect of nonequilibrium spin ($\delta P_E = 0$)] were also considered by Flatté *et al.* (2003) and Lebedeva and Kuivalainen (2003).

The current amplification (gain) $\beta = I_C/I_B$ (see Fig. 32) is typically ~ 100 in practical transistors. This ratio depends on many factors, such as the doping densities, carrier lifetimes, diffusion coefficients, and structure geometry. In an MBT β also depends on the spin splitting $2q\zeta$ (see Fig. 32) and the nonequilibrium polarization δP_E . This additional dependence of β in an MBT is called magnetoamplification (Fabian and Žutić, 2004). An important prediction is that the nonequilibrium spin can be injected at low bias all the way from the emitter, through the base, to the collector (Fabian *et al.*, 2002a; Fabian and Žutić, 2004) in order to make possible an effective control of β by δP_E .

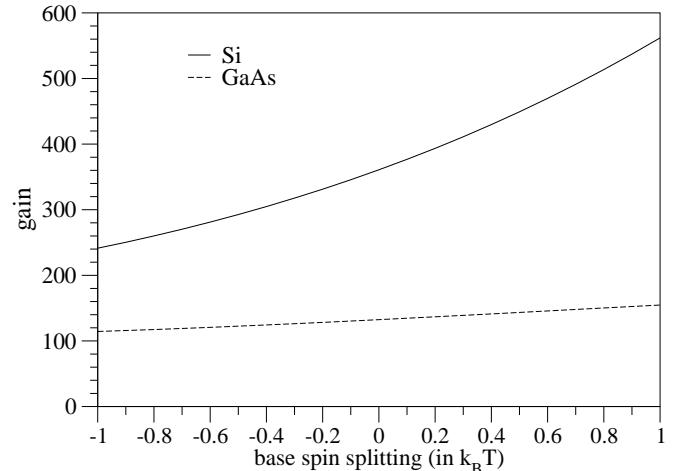


FIG. 33 Calculated gain dependence of an MBT as a function of base spin splitting $2q\zeta$, given in units of thermal energy $k_B T$. The nonequilibrium spin polarization in the emitter is $\delta P_E = 0.9$. Si (solid) and GaAs (dashed) materials parameters were applied. Adapted from Fabian *et al.*, 2002a.

The calculated dependence of the gain on the spin splitting for $\delta P_E = 0.9$ is shown in Fig. 33, for GaAs and Si materials parameters. The gain is very sensitive to the equilibrium magnetization in Si, while the rapid carrier recombination in GaAs prevents more effective control of the transport across the base. In Si it is the spin injection at the emitter-collector depletion layer which controls the current. As the spin-charge coupling is most effective across the depletion layer (see Sec. IV.D), this coupling is essential for the current in Si. In the limit of slow carrier recombination (Fabian *et al.*, 2002a),

$$\beta \sim \cosh(q\zeta/k_B T)(1 + \delta P_E P_{B0}). \quad (116)$$

Both magnetic field (through ζ) and nonequilibrium spin affect the gain, an implication of the spin-voltaic effect (Žutić and Fabian, 2003; Žutić *et al.*, 2002). The sensitivity of the current to spin can be used to measure the injected spin polarization. If no spin source is present ($\delta P_E = 0$), there is no spin-charge coupling in the space-charge regions, unless at least two regions are magnetic. The only remaining effects on the $I - V$ characteristics come from the sensitivity of the carrier densities in the equilibrium magnetic regions to ζ [see Eq. (116) for the case of $\delta P_E = 0$].

The MBT is, in effect, a magnetic heterostructure transistor, since its functionality depends on tunability of the band structure of the emitter, base, or collector. The advantage of MBT, however, is that the band structure is not built-in, but can be tuned *during* the device operation by magnetic field or spin injection signals. The challenges to demonstrate the predicted phenomena in MBT are similar to those of magnetic bipolar diodes, see Sec. IV.D.

3. Hot-electron spin transistors

Spin transistors that rely on transport of hot (non-thermalized) carriers have the potential to serve of several different purposes. On the one hand, they could be used as a diagnostic tool to characterize spin- and energy-dependent interfacial properties, scattering processes, and electronic structure, relevant to spintronic devices.¹²⁰ On the other hand, hot-electron transistors are also of interest for their ability to sense magnetic fields, their possible memory applications, and a their potential as a source of ballistic hot-electron spin injection. Below we discuss two representative examples, a spin-valve transistor and a magnetic tunneling transistor.

The spin-valve or Monsma transistor provided an early demonstration of a hot-electron spin transistor and realization of a hybrid spintronic device that integrates metallic ferromagnets and semiconductors (Monsma *et al.*, 1995, 1998). A three terminal structure¹²¹ consisted of a metallic base (B) made of a ferromagnetic multilayers in a CPP geometry [as depicted in Fig. 3(a)] surrounded by a silicon emitter (E) and collector (C) with two Schottky contacts, formed at E/B and B/C interfaces.¹²² Forward bias V_{EB} controls the emitter current I_E of spin-unpolarized electrons, which are injected into a base region as hot carriers. The scattering processes in the base, together with the reverse bias V_{BC} , influence how many of the injected electrons can overcome the B/C Schottky barrier and contribute to the collector current I_C . Similar to the physics of GMR structures (Gijs and Bauer, 1997; Levy and Mertig, 2002) scattering in the base region strongly depends on the relative orientation of the magnetizations in the ferromagnetic layers. Electrons with spin which has magnetic moment opposite (antiparallel) to the magnetization of a ferromagnetic layer typically are scattered more than electrons with parallel magnetic moments, resulting in a spin-filtering

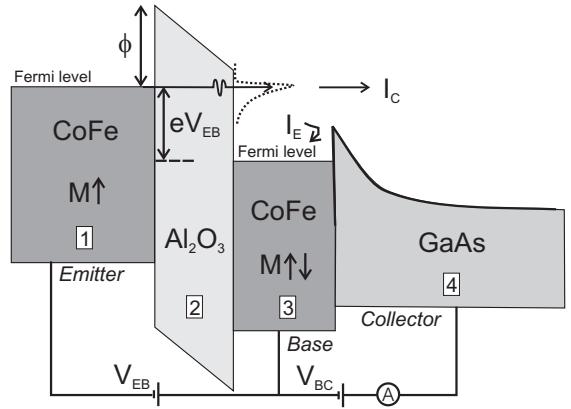


FIG. 34 Schematic energy diagram of a magnetic tunneling transistor. Region 1 is the emitter, region 2 the Al_2O_3 tunnel barrier of height ϕ , and region 3 the base. Together they form a magnetic tunnel junction. Region 4 is a semiconductor collector that has a Schottky barrier at the interface with the base. From van Dijken *et al.*, 2003c.

effect which can be described in terms of spin-dependent mean free path (Hong and Mills, 2000; Pappas *et al.*, 1991; Rendell and Penn, 1980). Generally, both elastic and inelastic scattering processes determine the effective spin-dependent mean free path, sometimes also referred to as the attenuation length.¹²³ The magnetoresistive response is usually expressed using magneto current (MC), defined as the change in collector current, normalized to the minimum value

$$MC = (I_{C\uparrow\uparrow} - I_{C\uparrow\downarrow})/I_{C\uparrow\uparrow}, \quad (117)$$

analogous to the expression for GMR or TMR structures [recall Eq. (1)], where $\uparrow\uparrow$ (parallel) and $\uparrow\downarrow$ (antiparallel) denote the relative orientation of the magnetizations. The large values of MC¹²⁴ ($>200\%$) and the sensitivity of $\sim 130\%$ per G measured at room temperature (Anil Kumar *et al.*, 2000) demonstrate a capability for magnetic-field sensors. Several important challenges, raised by the operation of the spin-valve transistor, need to be addressed to better realize the potential of hot-electron transistors. These challenges include increasing the small collector current and determining whether the spin injection of hot carriers into semiconductors is feasible. Furthermore, it would be desirable to fabricate structures in which semiconductor regions played an active role, not limited to energy selection (via Schottky

¹²⁰ These efforts are motivated in part by the success of (spin-insensitive) ballistic-electron-emission microscopy in providing high spatial and energy resolution of properties of metal/semiconductor interfaces (Bonnell Ed., 2001; Kaiser and Bell, 1988; Smith *et al.*, 2000). A subsequent variation—a ballistic-electron-magnetic microscopy, which also uses an STM tip to inject hot carriers, is capable of resolving magnetic features at a ~ 10 nm length scale (Rippard and Buhrman, 1999, 2000).

¹²¹ Similar to other hot-electron spin devices, the term transistor characterizes their three-terminal structure rather than the usual functionality of a conventional semiconductor transistor. In particular, a semiconductor bipolar transistor, which also has an emitter/base/collector structure, typically has a sizable current gain—a small change in the base current leads to a large change in the collector current (see Sec. IV.E.2). However, only a small current gain ~ 2 (due to large current in a metal base) was predicted in magnetic tunnel-junction-based devices (Hehn *et al.*, 2002).

¹²² Another realization of a spin-valve transistor combines a GaAs emitter with a Si collector (Dessein *et al.*, 2000).

¹²³ For electrons with sufficiently high excess energy, a scattering process (influencing the mean free path) does not necessarily remove the electron from the collector current. The attenuation length, which can be determined by measuring the base layer thickness dependence of the collector current [see (Rippard and Buhrman, 1999, 2000; van Dijken *et al.*, 2002b; Vlutters *et al.*, 2001)] can therefore differ from the effective mean free path.

¹²⁴ These values substantially exceed the CPP GMR value for the same magnetic multilayer used in the base.

barriers) of the carriers injected into the base and collector regions.

An alternative class of hot-electron transistors, often referred to as magnetic tunneling transistors, has a tunneling junction instead of a Schottky barrier emitter (Jiang *et al.*, 2003; Mizushima *et al.*, 1997; Sato and Mizushima, 2001; van Dijken *et al.*, 2002a,b, 2003c; Yamada and Mizushima, 1998). The addition of a tunnel junction, combined with a variable V_{EB} , allows exploration of hot-electron transport over an energy range of several eV. At large V_{EB} bias, the ratio I_C/I_E , important for the device performance, can be substantially increased over that of the spin-valve transistor (Sato and Mizushima, 2001; van Dijken *et al.*, 2003a,b).

A particular realization is depicted in Fig. 34. Different coercive fields in the regions 1 and 3 ensure independent switching of the corresponding magnetizations in the applied magnetic field. The magnetocurrent MC, defined in Eq. (117), shows a nonmonotonic behavior with V_{EB} (van Dijken *et al.*, 2003c) influenced by the conduction-band structure of a collector. In GaAs, in addition to the direct conduction band minimum at the Γ point [recall Fig. 6 (a)], there are indirect minima at L points at higher energy (Blakemore, 1982). After an initial decrease of MC with electron energy, at $V_{EB} \approx 0.3$ V larger than the base/collector Schottky barrier there is an onset of hot-electron transport into L valleys accompanied by an increase in MC (van Dijken *et al.*, 2003c).

A large magnetocurrent alone, measured in various hot-electron spin transistors (Monsma *et al.*, 1995, 1998; Sato and Mizushima, 2001; van Dijken *et al.*, 2003b), is not sufficient to demonstrate spin injection in a semiconductor collector. For conclusive evidence spin detection in a collector region is needed. This was first achieved (Jiang *et al.*, 2003) using optical detection with a spin LED structure¹²⁵ added to the collector in Fig. 34. Measurements at $T = 1.4$ K and $B = 2.5$ T, after a background subtraction, showed majority spin injection with $P_{\text{circ}} \approx 10\%$.

In another realization of a magnetic tunnel transistor, more similar to the original spin-valve transistor, the emitter was nonmagnetic (Cu) while the base was a magnetic multilayer (F1/N/F2) (van Dijken *et al.*, 2003b). The resulting strong spin-filtering effect can be inferred from the transmitted hot carriers with a spin-dependent exponential decay within the F_i , $i = 1, 2$ layer. Unpolarized electrons, injected from the emitter, after passing an F1 layer of thickness t acquire an effective transmitted polarization

$$P_{N1} = \frac{N_\uparrow - N_\downarrow}{N_\uparrow + N_\downarrow} = \frac{e^{-t/l_\uparrow} - e^{-t/l_\downarrow}}{e^{-t/l_\uparrow} + e^{-t/l_\downarrow}}, \quad (118)$$

where N_\uparrow and N_\downarrow represent the number of transmitted electrons with majority or minority spin and l_\uparrow and l_\downarrow are the corresponding attenuation length (the polarization P_{N2} has an analogous form). The resulting magnetocurrent can be expressed as (van Dijken *et al.*, 2003b)

$$MC = 2P_{N1}P_{N2}/(1 - P_{N1}P_{N2}), \quad (119)$$

which is analogous to Eq. (2) for TMR using Jullière's model, but with the redefined definition of spin polarization. At $V_{EB} = 0.8$ V and at $T = 77$ K, the measured MC exceeds 3400 %, while with Eq. (119) the polarization of the transmitted electrons can be estimated to exceed 90 %, even with a ferromagnet only a ~ 3 nm thick (van Dijken *et al.*, 2003b). A theoretical analysis of spin injection and spin filtering in magnetic tunneling transistors was given by Rashba (2003a) who extended the approach for ballistic spin injection (Kravchenko and Rashba, 2003) (Sec. II.C.2) to include the effects of hot-electron transport and inelastic scattering.

Future studies of hot-electron spin transistors are expected to result in increased spin injection even at room temperatures and to utilize other semiconductor collectors. It would be particularly desirable to demonstrate hot-electron spin injection in Si and facilitate an integration with the CMOS technology.

F. Spin qubits in semiconductor nanostructures

A potentially revolutionary idea in spintronics is the possibility of using the two-level nature of electron spin to create a solid-state quantum computer (Das Sarma *et al.*, 2001; DiVincenzo, 1995; Nielsen and Chuang, 2000). The basic unit in a quantum computer is the quantum bit (or qubit), the quantum analog of the binary bit in a classical digital computer. A qubit is essentially a controllable quantum two-level system (Das Sarma, 2001; Nielsen and Chuang, 2000). While the dimensionality (2^n) of the Hilbert space of n electron spins is the same as the number of configurations of a corresponding classical system, a quantum system can be in a superposition of all the basis states, effectively performing (via a unitary evolution) many classical computations in parallel. Several spin-based quantum computer schemes have been proposed and extensively studied¹²⁶ A common theme in these proposals is the idea of manipulating the dynamics of a single (or a few) electron spin(s) in semiconductor nanostructures (e.g., quantum dots), with the reasonable hope that the predicted behavior will extend to many-spin systems, requisite for practical quantum computation.

¹²⁵ Analogous to the spin LED from Fig. 12, in which GaAs collector served as an n -type spin aligner and InGaAs/GaAs was used for a quantum well.

¹²⁶ See, for example, (Burkard *et al.*, 1999; DiVincenzo, 2000; Friesen *et al.*, 2003; Hu and Das Sarma, 2000, 2001; Kane, 1998, 2000; Koiller *et al.*, 2002, 2003; Levy, 2002; Loss and DiVincenzo, 1998; Meier *et al.*, 2003; Piermarocchi *et al.*, 2002; Privman *et al.*, 1998; Skinner *et al.*, 2003; Troiani *et al.*, 2003; Vrijen *et al.*, 2000).

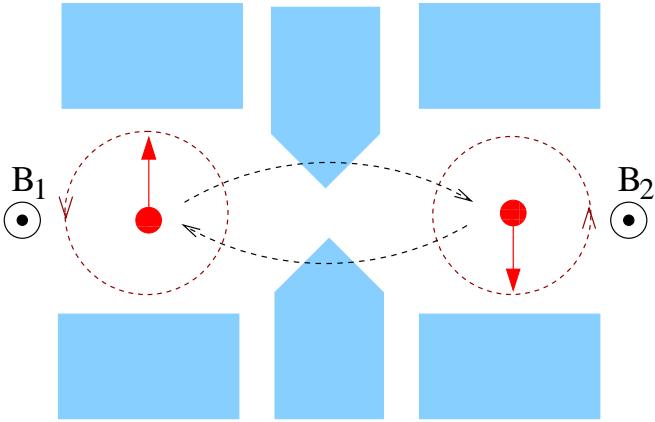


FIG. 35 The Loss and DiVincenzo (1998) proposal for spin-based solid-state quantum computing. Electrons are localized in electrostatically defined quantum dots, with coupling between electron spins—via the exchange interaction—allowed by tunneling between the dots. The tunneling is controlled by gate voltage. The figure shows two electrons localized in the regions defined by the gates (shaded). Single-qubit operations are realized by single-spin precessions (circles), performed by applying local magnetic fields (here perpendicular to the page) to each dot. Two-qubit operations are done through the exchange interaction indicated by the dashed curves. The scheme works according to the time-dependent Hamiltonian $H(t) = \sum'_{i,j} J_{i,j}(t) \mathbf{S}_i \cdot \mathbf{S}_j + \mu_B g \sum \mathbf{B}_i(t) \mathbf{S}_i$, where the first summation, over all neighboring spin pairs, describes the local exchange interaction (J is the exchange coupling), while the second describes the single spin operations by local magnetic fields. Variations to this scheme are described by Burkard *et al.* (2000).

The control of spin dynamics and entanglement [many-spin quantum correlations (Nielsen and Chuang, 2000)] at the single-spin level in a semiconductor quantum dot structure is a formidable task, which has not been achieved even at mK temperatures, although impressive experimental advances have recently been made (Elzerman *et al.*, 2003; Fujisawa *et al.*, 2002; Hanson *et al.*, 2003b). The current architectures for spin-based quantum computing employ GaAs quantum dots (Loss and DiVincenzo, 1998) or Si (or Si-Ge) systems (Kane, 1998), with different variations. The basic idea (see Fig. 35) is to manipulate the spin states of a single electron using external magnetic fields (or microwaves) for single-qubit operations and to utilize the quantum exchange coupling between two neighboring electrons to carry out two-qubit operations.

State-of-the-art techniques, to measure a single spin in a solid, such as magnetic resonance force microscopy (Barbic, 2002; Mamin *et al.*, 2003; Sidles *et al.*, 1995) or the spin-selective single-electron-transistor spectroscopy (Ono *et al.*, 2002), are still not sensitive enough for quantum computing operations. However, recently a single shot readout of the spin of an individual electron has been demonstrated using an electrical pump-probe measurement (Kouwenhoven, 2004). A single electron with

an unknown spin was trapped in a quantum dot for a few milliseconds. At the end of the trapping time the spin was measured by quickly shifting the Zeeman resolved spin states towards the Fermi energy. A spin to charge conversion allowed for an electrical readout of the spin.

The real motivation for using the spin as a qubit is its long coherence time, microseconds or longer in operational experimental conditions (de Sousa and Das Sarma, 2003a), to be contrasted with typical picosecond coherence times for charge or orbital states in solids. Interest in spin-based quantum computing will only increase as we understand more about spin coherence and relaxation from other spintronic studies. The broad subject of spin-based quantum computing, which is related to the areas of quantum measurement and quantum decoherence (Zurek, 2003) is beyond the scope of this review.

V. OUTLOOK

We have reviewed selected topics on spintronics, emphasizing both the fundamental aspects of spin dynamics, transport, and relaxation, and the potential applications. While the current push for spintronics is driven by the prospect of technological applications, the fundamental spin physics, which has a longstanding tradition in the solid-state community, is by itself exciting and worth pursuing. Furthermore, even though many proposed spintronic device schemes may turn out to be impractical in the end, their importance lies in stimulating interesting experimental and theoretical research.

There are many challenges and open questions to be tackled by future research, in particular a robust spin injection into silicon.¹²⁷ While GaAs is of great technological importance, the control of spin in silicon would raise hopes for seamless integration of spintronics with the current information technology. In addition, the small magnitude of the spin-orbit interaction and the absence of inversion symmetry lead to relatively long room-temperature spin lifetimes (of about 10 ns; see Sec. III.D.1), relaxing some constraints on the operational length and time scales. Important materials advances have been realized in improving the compatibility of Si/III-V structures (Sieg *et al.*, 1998) suggesting a possibility that the existing control of spin in GaAs or in III-V ferromagnetic semiconductors might be extended to Si.

Future progress in spin-polarized transport will be largely driven by the materials advances. In the context of semiconductors, considering all-semiconductor structures rather than the hybrid structures with metallic ferromagnets will depend on the improvements in fer-

¹²⁷ Small signals attributed to spin injection have already been reported (Jia *et al.*, 1996).

romagnetic semiconductors, for example, whether they can achieve higher mobility, higher Curie temperature,¹²⁸ and a simple fabrication of high quality interfaces with nonmagnetic materials. What is missing, even in the currently available materials, is a systematic understanding of the effects of magnetic interfaces and materials inhomogeneities on spin-polarized transport. A comprehensive transport calculation in the actual devices with realistic electronic structure of the studied materials would provide valuable insights into both the spin polarization being measured and how it is reduced from the moment it was generated.

Spin relaxation and spin dephasing of conduction electrons is a rather field, with the basic principles well understood. What is needed are accurate band-structure-derived calculations of spin relaxation times, in both metals and semiconductors. The same can be said of *g* factor, calculation of which from first principles is a nontrivial task that has not been accomplished even for the elemental metals. An important and still debated issue is spin relaxation and decoherence of localized or confined electrons, where the hyperfine-interaction mechanism dominates. Furthermore, single-spin relaxation and decoherence, and their relation to the ensemble spin dephasing, need to be pursued further in the context of quantum computing. A first step towards understanding single-spin relaxation is the recent experiment of Hanson *et al.* (2003b) in a one-electron quantum dot.

While dynamic nuclear polarization induced by electron spin can often be a nuisance for detecting intrinsic spin dynamics (see Sec. III.D.3), the interaction between electron and nuclear spins (Fleisher and Merkulov, 1984a; Paget and Berkovits, 1984; Smet *et al.*, 2002; Vagner, 2003) is of fundamental importance for spintronics. An NMR of the nuclear spin polarized by spin-polarized photoexcited electrons has already been used to detect the nonequilibrium electron spin in Si (Lampel, 1968). On the other hand, an NMR signal can be detected optically through measuring changes in the circular polarization of photoluminescence due to resonant variations of the nuclear field (D'yakonov and Perel', 1984a), as shown first in p-doped Ga_{0.7}Al_{0.3}As (Ekimov and Safarov, 1972). The early work of Lampel (1968), and Ekimov and Safarov (1972) established the basic principles for a series of experiments that demonstrated various realizations of an all-optical NMR. The role of the resonant radio waves is played by periodically optically ex-

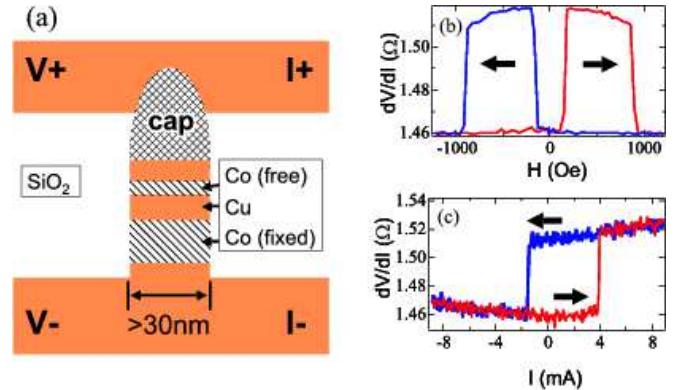


FIG. 36 (a) Nanopolar device: schematic diagram of a nanopolar device operating at room temperature. The direction of magnetization is fixed (pinned) in the thick Co film and free in the thin Co film; (b) differential resistance dV/dI of a nanopolar device as a function of applied field; (c) dV/dI of the device as a function of applied current. The arrows in panels (b) and (c) represent the direction of magnetic field and current sweeps, respectively. For positive current, electrons flow from the thin to the thick Co film. Adapted from Albert *et al.*, 2002.

cited electron spins (Eickhoff *et al.*, 2002; Fleisher and Merkulov, 1984b; Klevich, 1986; Klevich *et al.*, 1980, 1981; Kikkawa and Awschalom, 2000; Salis *et al.*, 2001a). Electron-nuclear spintronics is likely to become relevant for quantum computation and for few-spin manipulations, which can benefit from long nuclear spin coherence times (even lasting minutes).

The range of potential spintronic applications goes beyond the use of large magnetoresistive effects. Rudolph *et al.* (2003), for example, have demonstrated the operation of a spin laser. The laser is a vertical-cavity surface-emitting laser (VCSEL), optically pumped in the gain medium, here two InGaAs quantum wells, with 50% spin-polarized electrons. The electrons recombine with heavy holes, which are effectively unpolarized, emitting circularly polarized light (see Sec. II.B). The threshold electrical current, extracted from the pump power for the lasing operation, was found to be $0.5 \text{ A}\cdot\text{cm}^{-2}$, which is 23% below the threshold current of the spin-unpolarized VCSEL. Furthermore, for a fixed pump power, the emission power of the laser changed by 400% upon changing the degree of circular polarization of the pump laser. The reason for the decrease in threshold is the selective coupling of spin-polarized electrons to photons with one helicity. While the experiment was conducted at 6 K, a room-temperature operation and an electrically pumped laser should be viable as well.¹²⁹

The demonstration that the flow of spin-polarized car-

¹²⁸ There still remain many challenges in accurately predicting Curie temperature. First principles results suggest that dominant models of ferromagnetism in semiconductors cannot be used to explain a variation of Curie temperature across different materials (Erwin and Žutić, 2004). For reviews of ferromagnetic semiconductor theories outside the scope of this article, see, for example, Nagaev (1983); Bhat *et al.* (2002); Dietl (2002); Sanvitto *et al.* (2002), Das Sarma *et al.* (2003); König *et al.* (2003); Timm (2003).

¹²⁹ The requirement is that the spin relaxation time be longer than the carrier recombination time, and that the spin injection spot and the gain medium be within the spin transport length.

riers, rather than applied magnetic field, can also be used to manipulate magnetization of ferromagnetic materials brings the exciting prospect of a novel class of spintronic devices. In addition to reversal of magnetization, which is a key element in realizing various magnetoresistive applications, the driving of a spin-polarized current can lead to coherent microwave oscillations in nanomagnets (Kiselev *et al.*, 2003). Spin-transfer torque (Sec. I.B.1) has already been realized in several experimental geometries. These include nanowires (Kelly *et al.*, 2003; Wengrove *et al.*, 1999), point contacts (Ji *et al.*, 2003; Tsui *et al.*, 1998, 2000, 2002), nanoconstrictions (Myers *et al.*, 1999; Rippard *et al.*, 2004), and nanopillars (Albert *et al.*, 2002; Katine *et al.*, 2000; Urazhdin *et al.*, 2003) (see Fig. 36), all involving metallic ferromagnets. The common feature of all these geometries is a need for very large current densities ($\sim 10^7 \text{ Acm}^{-2}$). Ongoing experiments (Chiba *et al.*, 2004; Munekata, 2003; Yamanouchi *et al.*, 2004) to demonstrate spin-transfer torque (together with other cooperative phenomena) in ferromagnetic semiconductors, which have much smaller magnetization than their metallic counterparts, are expected to also require much smaller switching currents. Based on the findings in electric-field controlled ferromagnetism (see Fig. 27), it has been demonstrated that the reversal of magnetization in (In,Mn)As can be manipulated by modifying the carrier density, using a gate voltage in a FET structure (Chiba *et al.*, 2003).

Acknowledgments

We thank R. H. Buhrman, K. Bussmann, R. de Sousa, M. I. D'yakonov, S. C. Erwin, M. E. Fisher, A. M. Goldman, K. Halterman, X. Hu, S. V. Iordanskii, M. Johnson, B. T. Jonker, K. Kavokin, J. M. Kikkawa, S. Maekawa, C. M. Marcus, I. I. Mazin, B. D. McCombe, J. S. Moodera, H. Munekata, B. E. Nadgorny, H. Ohno, S. S. P. Parkin, D. C. Ralph, E. I. Rashba, W. H. Rippard, V. I. Safarov, G. Schmidt, E. Sherman, R. H. Silsbee, D. D. Smith, M. D. Stiles, O. T. Valls, S. van Dijken, H. M. van Driel, T. Venkatesan, S. von Molnár, and J. Y. T. Wei for useful discussions. We thank D. D. Awschalom, N. C. Emley, R. T. Harley, M. Johnson, B. T. Jonker, K. Kavokin, J. M. Kikkawa, C. M. Marcus, H. Munekata, Y. Ohno, A. Oiwa, S. S. P. Parkin, R. J. Soulen, Jr., and M. Tanaka for providing us with illustrative figures from their published works, and A. Kaminski and D. J. Priour for help with preparation of the manuscript. This work was supported by DARPA, the US ONR, and the NSF-ECS.

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2DEG	two dimensional electron gas
BAP	Bir-Aronov-Pikus
BCS	Bardeen Cooper Schrieffer
BIA	bulk inversion asymmetry
BTK	Blonder-Tinkham-Klapwijk
CESR	conduction electron spin resonance
CIP	current in plane
CMOS	complementary metal oxide semiconductor
CMR	colossal magnetoresistance
CPP	current perpendicular to plane
DNA	deoxyribonucleic acid
DP	D'yakonov-Perel'
EDSR	electron dipole spin resonance
EMF	electromotive force
ESR	electron spin resonance
EY	Elliot-Yafet
F	ferromagnet
FSm	ferromagnetic semiconductor
GMR	giant magnetoresistance
FET	field effect transistor
HFI	hyperfine interaction
I	insulator
HTSC	high temperature superconductor
LED	light emitting diode
MBD	magnetic bipolar diode
MBE	molecular beam epitaxy
MBT	magnetic bipolar transistor
MC	magnetocurrent
MR	magnetoresistance
MRAM	magnetic random access memory
MTJ	magnetic tunnel junction
N	normal (paramagnetic) metal
NMR	nuclear magnetic resonance
QD	quantum dot
QPC	quantum point contact
QW	quantum well
S	superconductor
SET	single electron transistor
SFET	spin field effect transistor
SIA	structure inversion asymmetry
Sm	semiconductor
SQUID	superconducting interference quantum device
STM	scanning tunneling microscope
TESR	transmission electron spin resonance

TABLE II List of acronyms used in the text.