

# Quantum transport in Rashba spin-orbit materials: A review

Dario Bercioux\*

*Donostia International Physics Center (DIPC),  
Manuel de Lardizbal 4,  
E-20018 San Sebastián,  
Spain*

*IKERBASQUE, Basque Foundation of Science,  
48011 Bilbao, Basque Country,  
Spain*

*Dahlem Center for Complex Quantum Systems and Institut für Theoretische Physik,  
Freie Universität Berlin,  
Arnimallee 14, D-14195 Berlin,  
Germany*

Procolo Lucignano†

*CNR-SPIN, Monte S. Angelo – via Cinthia, I-80126 Napoli,  
Italy  
Dipartimento di Fisica,  
Università di Napoli “Federico II”,  
Monte S. Angelo, I-80126 Napoli,  
Italy*

In this review article we describe spin-dependent transport in materials with spin-orbit interaction of Rashba type. We mainly focus on semiconductor heterostructures, however we consider topological insulators, graphene and hybrid structures involving superconductors as well. We start from the Rashba Hamiltonian in a two dimensional electron gas and then describe transport properties of two- and quasi-one-dimensional systems. The problem of spin current generation and interference effects in mesoscopic devices is described in detail. We address also the role of Rashba interaction on localisation effects in lattices with nontrivial topology, as well as on the Ahronov-Casher effect in ring structures. A brief section, in the end, describes also some related topics including the spin-Hall effect, the transition from weak localisation to weak anti localisation and the physics of Majorana Fermions in hybrid heterostructures involving Rashba materials in the presence of superconductivity.

## CONTENTS

I. Introduction	1	V. Interference effects and Berry phase	17
II. Materials with Rashba Spin-orbit interaction	3	A. Rashba interaction as SU(2) gauge field: application to quantum networks	17
A. Semiconductors heterostructures	3	B. Interference in quantum rings	19
B. Quantum wires	4	VI. Related problems	21
C. Carbon-based materials	4	A. Spin-Hall effect	21
D. Topological insulators	5	B. Diffusive limit: weak anti-localization	22
E. Weyl semimetals	6	C. Quantum wires with RSOI, superconductivity and magnetic field: The quest for Majorana Fermions	23
III. Spin transport in RSOI material	7	VII. Conclusions and outlook	25
A. General properties of the RSOI in semiconductors	7	Acknowledgments	26
B. Rashba SOI in quantum wires: subbands hybridisation	8	References	26
C. General properties of the Rashba SOI in graphene	9		
D. The spin double refraction	10		
IV. Pure spin current generation in RSOI material	11		
A. Pure spin current generators	13		
1. Spin Ratchet	13		
2. Spin Pumping	14		
B. Datta and Das spin Field Effect Transistor	14		

## I. INTRODUCTION

All the modern electronic devices are based on a classical paradigm of negative carriers — electrons — and positive charge carriers — holes. This leaves completely unused an additional degree-of-freedom carried by both carriers: the spin. Spintronics explores phenomena that interlink the spin and the charge degree-of-freedom. It

\* [dario.bercioux@dipc.org](mailto:dario.bercioux@dipc.org)

† [procolo.lucignano@spin.cnr.it](mailto:procolo.lucignano@spin.cnr.it)

is the field where traditional solid-state physics and material research have created their strongest bond, with each taking alternate leading roles. From our perspective, spintronics can be mainly divided into two distinct subfields: metal and semiconductor spintronics (Žutić *et al.*, 2004; Sinova and Žutić, 2012).

Metallic spintronic devices originate from the discovery of giant magnetoresistance (GMR) in 1988 (Baibich *et al.*, 1988; Binasch *et al.*, 1989) and the subsequent development of the spin valve (Dieny *et al.*, 1991). The GMR effect can be understood by assuming that any spin current is carried by two "types" of carriers, spin-up and spin-down. The two-channel picture of spin transport proposed by Mott explains the behaviour of magnetoresistive devices (Camley and Barnaś, 1989), including GMR and tunnelling magnetoresistance (TMR) (Moodera *et al.*, 1995), as well as spin injection into metals (Johnson and Silsbee, 1988).

In the GMR effect, two ferromagnetic layers sandwich a non-ferromagnetic metal layer of nanometer thickness. When the magnetisation of the two ferromagnetic layers is parallel, the valve is in a low resistance state (open). When the two are antiparallel, the valve is in a high resistance state (closed). The TMR effect rather than the GMR one, takes place when a metallic spacer is substituted by an insulating barrier. In TMR at room temperature, the spin-valve effect increases by a factor of ten with respect to GMR. This is the basic principle of hard disk read heads and magnetic random access memories. These are some of the most successful technologies of the past decades, with scaling trends outdoing even complementary metal-oxide-semiconductor, the technology for constructing integrated circuits. Albert Fert and Peter Grünberg have been awarded the Physics Nobel Prize in 2007 for their studies on the GMR effect.

Semiconductor spintronics, on the other hand, has the potential to be integrated seamlessly with nowadays' semiconductor electronics. It is progressing along a similar path as metallic spintronics and has achieved a remarkable success in the past decade. The spin-field effect transistor (spin-FET) was proposed in 1990 (Datta and Das, 1990) as a development to the GMR set-up where the two ferromagnets are left collinear and the "on" and "off" status is achieved by rotating the injected spin when travelling through the two contacts. This geometry does not require a magnetic field for switching the relative magnetisation of the two ferromagnets, thus reducing spurious magnetic fields into the electronic circuits. The spin-FET is an electronic analog of an electro-optical modulator for photons. A ferromagnet injects spin polarised carriers into a semiconductor channel. Here, due to the modulation of the spin-precession length, it is possible to vary the spin orientation. This modulated spin signal is then detected by a second ferromagnet. The spin precession length is varied via top- and back-gate voltages that tune the strength of the spin-orbit inter-

action (SOI) of Rashba (R) type (Rashba, 1960; Yu A. Bychkov and Rashba, 1984). The RSOI originates from the lack of inversion symmetry along the growth direction of the semiconductor heterostructure that is hosting a two-dimensional electron gas (2DEG) (de Andrade e Silva *et al.*, 1994, 1997). It can be tuned by changing the shape of the confining potential via the application of an external electric field (Nitta *et al.*, 1997; Schäpers *et al.*, 1998; Grundler, 2000; Miller *et al.*, 2003; Meier *et al.*, 2007). However, the conductivity mismatch problem (Johnson and Silsbee, 1988; Schmidt *et al.*, 2000) between a ferromagnet and a low-dimensional semiconductor has hindered, for a long time, an efficient injection of spin polarised currents. Progress for solving this problem came with the research on ferromagnetic semiconductors (Ohno, 1998). Nevertheless, only recently it has been possible to implement completely the functionality of the spin-FET first in Ref. (Koo *et al.*, 2009), and after, with minor changes, in Ref. (Chuang *et al.*, 2014).

Semiconductor spintronics is also relevant because it has permitted to observe other coherent phenomena, such as the Aharonov-Casher (AC) effect (Aharonov and Casher, 1984) in ring-type and other loop structures (Yau *et al.*, 2002) so as proposed by many groups (Nitta *et al.*, 1999; Frustaglia and Richter, 2004; Bercieux *et al.*, 2005a; Capozza *et al.*, 2005; Lucignano *et al.*, 2007). The Aharonov-Casher effect is the analog of the Aharonov-Bohm (AB) effect (Aharonov and Bohm, 1959) for particles with an angular momentum in an external electrical field (Richter, 2012). In the case of carriers in a semiconductor 2DEG, the RSOI permits the coupling of the electron spin with the external electric field. Other phenomena, *e.g.*, the spin-Hall effect (D'yakonov and Perel', 1971; Hirsch, 1999; Murakami *et al.*, 2003; Sinova *et al.*, 2004), have been predicted and observed in semiconductors (Kato *et al.*, 2004; Sih *et al.*, 2005a; Wunderlich *et al.*, 2005a,b; Sih *et al.*, 2006a) before their observation in metals (Valenzuela and Tinkham, 2006). A key role in the success of semiconductor spintronics is the tunability of the RSOI via external gating.

The material in this review article is presented in the following form: in Sec. II we present a summary of the materials with sizable RSOI effects, these range from the standard semiconductor heterostructures, carbon-based materials, topological insulators (TIs) to Weyl semimetals (WSs). In Sec. III, in order to introduce some basic transport property of carriers in systems with RSOI, we revise the spectral properties of 2DEGs and graphene with RSOI, in addition we spend some effort to underline the basic difference in the presence of a further lateral confinement, thus moving from the two-dimensional (2D) to the quasi-one-dimensional-case (quasi-1D). After, we introduce the concept of spin-double refraction that is a phenomenon taking place in a hybrid structure composed of region with RSOI sandwiched between two regions without RSOI. In Sec. IV we give a defini-

tion of pure spin currents and propose a derivation of a formula for a spin current in the standard Landauer-Büttiker approach, furthermore we introduce two proposals for generating pure spin currents — based on the ratchet and the pumping effect. Next, we give an in-depth analysis of the spin-FET by Datta and Das and an explanation of its working principle, finally we present a recent experiment showing a possible realisation of such a transistor. In Sec. V we give a complementary look at the effects induced by RSOI, specifically we will analyse its effects as non-Abelian gauge field. Particularly, we look at non-Abelian effects in quantum networks and quantum rings. In Sec. VI we give a very brief and general overview over other aspects of the physics associated to the RSOI, specifically we look at the spin-Hall effect (SHE) in Sec. VI.A, than, in Sec. VI.B, we will see how RSOI can modify the weak localisation (WL) into weak anti-localisation (WAL) in a disordered system. Finally in Sec. VI.C we will quickly review the vibrant field of the quest of Majorana quasi particles in quantum wires with RSOI, magnetic field and superconductivity. In Sec. VII we present our outlook for this growing field of condensed matter physics.

## II. MATERIALS WITH RASHBA SPIN-ORBIT INTERACTION

On a very general basis, SOI routes down to the relativistic correction to hydrogen-like atoms (Bransden and Joachain, 2003; Sakurai and Napolitano, 2014). It reads

$$\begin{aligned} \mathcal{H}_{\text{SO}} &= -\frac{2\hbar}{(2mc)^2} \mathbf{s} \cdot (\mathbf{E}(\mathbf{r}) \times \mathbf{p}) \\ &= \frac{1}{r} \frac{dV(r)}{dr} \mathbf{s} \cdot (\mathbf{r} \times \mathbf{p}) \end{aligned} \quad (1)$$

where  $\mathbf{p}$  is the momentum operator and  $\mathbf{s} = (s_x, s_y, s_z)$  is a vector of Pauli matrices describing the spin  $\mathbf{S} = \frac{\hbar}{2} \mathbf{s}$  operator. By considering a Coulomb electrostatic potential  $V(r) = -\frac{Ze^2}{r}$  we have that the strength of this relativistic correction goes as  $\frac{Ze^2}{r^3}$ , thus proportional to the atomic number  $Z$  (Bransden and Joachain, 2003; Sakurai and Napolitano, 2014). From the relativistic correction expressed in Hamiltonian (1) we learn that materials characterised by a higher atomic number will present in general a stronger SOI, this will be the case for TIs and Weyl- and Dirac-semimetals.

### A. Semiconductors heterostructures

In this review we shall mostly discuss the physics of low-dimensional semiconductor heterostructures. The simplest example is a 2DEG. This is commonly realised in III-V semiconductor heterostructures by modulating

the doping density along the growth direction — usually (001) (Bastard, 1988; Seeger, 2004). In solids, SOI can have different nature depending on the crystal symmetries (Samokhin, 2009), however in this review article we shall mostly deal with the two main independent sources of SOI in III-V semiconductors, namely the RSOI, due to the lack of structural inversion symmetry (SIA) (de Andrada e Silva *et al.*, 1994, 1997) and a Dresselhaus (D) term due to a lack of bulk inversion symmetry (BIA) (Dresselhaus, 1955). The three-dimensional (3D) SOI correction to the free electronic Hamiltonian due to BIA reads:

$$\begin{aligned} \mathcal{H}_{\text{DSOI}}^{3D} &\propto p_x (p_y^2 - p_z^2) s_x + p_y (p_z^2 - p_x^2) s_y \\ &\quad + p_z (p_x^2 - p_y^2) s_z. \end{aligned} \quad (2)$$

In order to achieve the effective Hamiltonian acting on the electrons confined in the 2DEG we have to integrate along the growth direction with the constraint that  $\langle p_z \rangle = 0$ , while  $\langle p_z^2 \rangle \neq 0$  is a sample dependent constant. Therefore we have

$$\mathcal{H}_{\text{DSOI}} = \beta (p_y s_y - p_x s_x) + \gamma (p_x p_y^2 s_x - p_y p_x^2 s_y) \quad (3)$$

where  $\gamma$  is a material dependent constant, and  $\beta$  depends on  $\langle p_z^2 \rangle$ . The first term contains the so called linear DSOI whereas the second one describes the cubic one (Winkler, 2003). Usually the latter can be disregarded as  $\langle p_z^2 \rangle \ll p_x, p_y$  in a 2DEG.

Due to charging effects, the quantum well confining electrons in two dimensions is never perfectly flat, therefore at the interface between the two differently doped semiconductors, a potential gradient arises  $\nabla V = \mathbf{E}$ . This effective electric field couples with the electrons motion as

$$\mathcal{H}_{\text{RSOI}} \propto (\mathbf{E} \times \mathbf{p}) \cdot \mathbf{s}. \quad (4)$$

Considering a quantum well along the (001) growth direction  $\mathbf{E} = E_z \mathbf{z}$ , the RSOI is rephrased as:

$$\mathcal{H}_{\text{RSOI}} = \frac{\alpha}{\hbar} (\mathbf{s} \times \mathbf{p}) \cdot \mathbf{z} = \frac{\alpha}{\hbar} (p_y s_x - p_x s_y), \quad (5)$$

where  $\alpha$  depends on the material and on the confining potential. By simple inspection one can notice that Rashba Hamiltonian (5) and linear Dresselhaus SOI (3) are equivalent under the unitary transformation rotating the spin Pauli matrices  $s_x \rightarrow s_y$ ,  $s_y \rightarrow s_x$  and  $s_z \rightarrow -s_z$  therefore the spectral properties of electrons subject to RSOI or linear DSOI are exactly the same (Lucignano *et al.*, 2008). It is important to notice that through a simple mathematical manipulation we can recast RSOI in the form  $\mathbf{s} \cdot \mathbf{B}_{\text{eff}}(\mathbf{p})$  where  $\mathbf{B}_{\text{eff}} \propto (\mathbf{z} \times \mathbf{p})$  is an effective magnetic field rotating the carrier spin in the plane of the 2DEG.

Despite this formal equivalence, the role to RSOI has been investigated with much more attention in the last

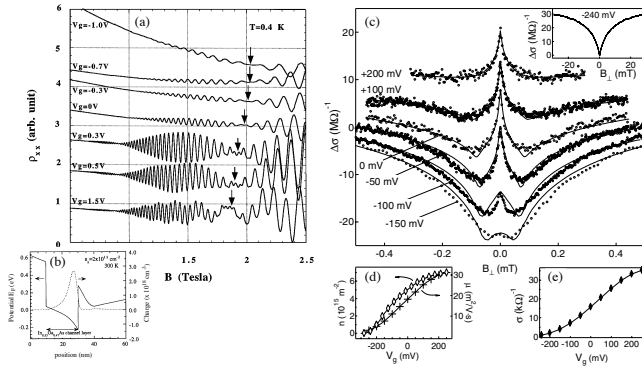


FIG. 1 (a) Schubnikov-de Haas oscillations as function of the gate voltages (Nitta *et al.*, 1997); (b) band diagram and electron distribution. Panels (a) and (b) courtesy of Ref. [Nitta *et al.*, 1997]; (c) Experimental magneto-conductance,  $\Delta\sigma = \sigma(b) - \sigma(0)$  (circles), offset for clarity, along with three-parameter fits (solid line) for several gate voltages. Inset: Experimental magneto-conductance data for the most negative gate voltage, showing pure weak localization. (d) Density and mobility as function of  $V_g$ , extracted from longitudinal and Hall voltage measurements; (e) Experimental conductivity, showing strong dependence on  $V_g$ . Panels (c)-(e) courtesy of Ref. [Miller *et al.*, 2003].

decades for two main reasons. First, in conventional III-V 2DEGs  $\alpha > \beta$ . Second, more interestingly, from few decades  $\alpha$  can be easily tuned by gating the heterostructure (de Andrada e Silva *et al.*, 1994, 1997) to some hundreds of meV nm. The spin precession length can be expressed as

$$l_{SO} = \frac{\pi}{k_{SO}} = \frac{\pi\hbar^2}{\alpha m^*},$$

in III-V quantum wells it ranges from hundreds of nm to few  $\mu m$ , thus comparable with typical dimensions achievable in the fabrication of modern mesoscopic devices. The RSOI was first predicted in bulk semiconductors by Rashba (Rashba, 1960). Later Bychkov and Rashba (Yu A. Bychkov and Rashba, 1984) proposed to use the Hamiltonian (5) in order to explain some magnetic properties of semiconductor heterostructures (Stein *et al.*, 1983; Stormer *et al.*, 1983). In figure 1(b) we show the shape of the quantum well and of the electron density profile showing the position of the 2DEG. The lack of inversion symmetry in semiconducting 2DEG heterostructures is at the origin of the RSOI. Therefore a change in the asymmetry can also lead to a change in the strength of the RSOI. For example, by applying a gate voltage is possible to modify the profile of the potential confining the 2DEG with a secondary effect of changing the strength of the RSOI. Thus, in a first approximation the coupling constant  $\alpha$  depends on the gate voltage  $\alpha \rightarrow \alpha(V)$ .

A change in the strength of the  $\alpha$ , thus, can produce a change in the beating pattern of the Schubnikov-de

Haas oscillations (de Andrada e Silva *et al.*, 1994, 1997) [c.f. Figure 1(a)]. This effect was first observed by the group of Nitta (Nitta *et al.*, 1997) and after by other groups (Schäpers *et al.*, 1998; Grundler, 2000). Another method for probing the variation of the coupling constant is to observe the transition from WL to weak WAL (Miller *et al.*, 2003; Akkermans and Montambaux, 2007)<sup>1</sup>, see for instance figure 1(c)-(e).

## B. Quantum wires

By applying a further confinement to a 2DEG it is possible to confine the electrons in a quasi-1D structure — a quantum wire. A detailed investigation of the RSOI in InGaAs/InP quantum wires of different width has been put forward by Schäpers *et al.* [Schäpers *et al.*, 2004, 2006]. Quantum wires can also be made out of InAs, GaAs, InSb or other materials (Nilsson *et al.*, 2009; Nadj-Perge *et al.*, 2012; Estévez Hernández *et al.*, 2010). Recently, quantum wires with sizable RSOI are attracting a lot of interest as they are the fundamental building block for many proposals aiming at the observation of Majorana quasi particles in condensed matter physics. In these setups the interplay between RSOI, superconductivity and magnetic fields (Oreg *et al.*, 2010; Lutchyn *et al.*, 2010) is pivotal.<sup>2</sup>

## C. Carbon-based materials

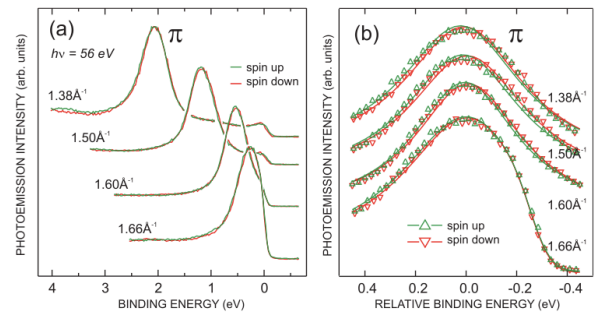


FIG. 2 (Colours online) Spin-resolved photoemission spectra along  $\overline{\Gamma K}$  in the vicinity of the Fermi energy. (a) Overview spectra and (b)  $\pi$  states at arbitrary offset. Courtesy of Ref. [Varykhalov *et al.*, 2008].

<sup>1</sup> More details about this phenomenon will be given in Sec. VI.B.  
<sup>2</sup> Some more analysis of Majorana physics is reported in Sec. VI.C.

The RSOI has been predicted also for single layer graphene (SLG), using simple symmetry arguments (Kane and Mele, 2005a,b). A more rigorous calculation has been proposed by Huertas-Hernando *et al.* (Huertas-Hernando *et al.*, 2006), where a modulation via atomic Stark effect and curvature effects were considered. However, due to the small atomic number of the carbon atoms [see equation (1)] the strength of the coupling  $\alpha$  is very small in the SLG structure. There are proposal for overcoming this obstacle and increasing the strength of the SOI, by coating the graphene surface with *ad*-atoms thus inducing a spin-dependent hopping mediated by the surface impurities. This is the case, for example, of SLG deposited on a Ni substrate and intercalated with Au atoms. Spin-resolved angle resolved photoemission spectroscopy measurements have shown a sizable RSOI of the order of some meV (Varykhalov *et al.*, 2008; Dedkov *et al.*, 2008; Li *et al.*, 2011; Marchenko *et al.*, 2012). Similar proposal have been advanced also for hydrogen impurities (Castro Neto and Guinea, 2009). The modification to the energy spectrum of SLG due to RSOI are shown in figure 2. Large RSOI has been also proposed by using rotating magnetic fields (Klinovaja and Loss, 2013).

The RSOI in SLG is described by a Hamiltonian very similar to (5), the main difference is that it depends only on the pseudo-spin and not on the momentum operators. To the lowest order in the momentum expansion:

$$\mathcal{H}_{\text{SO}}^{\text{SLG}} = \frac{\lambda}{2} (\mathbf{s} \times \boldsymbol{\sigma})_z, \quad (6)$$

where  $\mathbf{s}$  are the Pauli matrices describing the electron spin and  $\boldsymbol{\sigma}$  its pseudo-spin. Here the pseudo-spin is the internal degree of freedom describing the presence of two inequivalent carbon atoms in the honeycomb lattice of SLG (Neto *et al.*, 2009).

The RSOI has been predicted also in bi-layer graphene, due to the interaction with a substrate of and an external electric field (Gelderen and Smith, 2010; Guinea, 2010; Kunschuh *et al.*, 2012; Mireles and Schliemann, 2012), however so far it is has not been measured.

In order to conclude this overview of carbon-based materials with RSOI, relevant effects are predicted also for carbon nanotubes (CNTs) (Martino *et al.*, 2002; Martino and Egger, 2005; Izumida *et al.*, 2009). In Refs. [Kuemeth *et al.*, 2008 and Jespersen *et al.*, 2011] the RSOI interaction strength is measured in CNT quantum dots.

#### D. Topological insulators

Topological insulators are band insulators hosting spin polarised edges states within the bulk gap (Hasan and Kane, 2010; Qi and Zhang, 2011; Ando, 2013).

The original idea about this new state of matter was put forward by Kane and Mele in two seminal research

papers published in 2005 (Kane and Mele, 2005a,b). They showed that a specific type of SOI in SLG can lead to the opening of a gap in the spectrum and that edge states exists within this gap when considering a ribbon geometry. In figure 3 we can observe the gap opened by the intrinsic SOI (Huertas-Hernando *et al.*, 2006) and the edge states within the gap. In two-dimensions, this phenomenon is also known as quantum spin-Hall effect (QSHE). It can be thought as two copies (one per spin) of the integer quantum Hall effect (IQHE). The main difference is that, contrary to the magnetic field, SOI does not break time-reversal symmetry. Thus, here, there are two counter propagating helical edge states opposed to the single chiral edge state of the quantum Hall fluid. As in the case of the IQHE, the edge states are characteristics only of the finite size systems. It is also possible to introduce a specific topological numbers named  $\mathbb{Z}_2$  that defines the robustness of these edge states (Kane and Mele, 2005a).

Up to now, such phenomenon has not been observed in SLG, however shortly after the seminal works of Kane and Mele, a proposal for observing the QSHE in HgTe/CdHgTe quantum wells was put forward by Bernevig *et al.* [Bernevig *et al.*, 2006 and Bernevig and Zhang, 2006], whose prediction was readily confirmed by König *et al.* [König *et al.*, 2006]. Since than, there is a huge quest for discovering new 2D TIs. One of the most interesting is represented by InAs/GaSb quantum well (Liu *et al.*, 2008; Knez *et al.*, 2011) which is very attractive as it is achievable using the standard and, very well known, III-V semiconductor technology.

Contrary to the case of IQHE, the paradigm of the QSHE can be extended to 3D. This work has been mainly carried out by Fu and Kane [Fu *et al.*, 2007 and Fu and Kane, 2007]. Since their seminal works, a race for discovering new 3D TIs is open. Recent overviews on these materials can be found in one of the several review articles that has been published on the topic (Hasan and Kane, 2010; Qi and Zhang, 2011; Ando, 2013). Among them the paper by Ando (Ando, 2013) contains an interesting table summarising all known 2D and 3D TI discovered up to 2013.

Most of the research work is devoted to analyse the spectral properties of such material—mainly via angle-resolved photoemission spectroscopy. Only few experiments show the transport properties of this new class of materials. Mainly because most of the 3D TI are not proper insulators due to a non negligible current flowing through impurity states (Bardarson and Moore, 2013). A very interesting set of transport experiments has been performed in films of Bi<sub>2</sub>Se<sub>3</sub> (Qu *et al.*, 2011) prepared with a special two-dimensional geometry (c.f. figure 4). In this work the authors show a sizable tunability of RSOI in 3D TI. The effects of this tunability are clearly observed in the interference pattern of the Aharonov-Bohm (AB), the Altshuler-Aronov-Spivak (AAS) and the

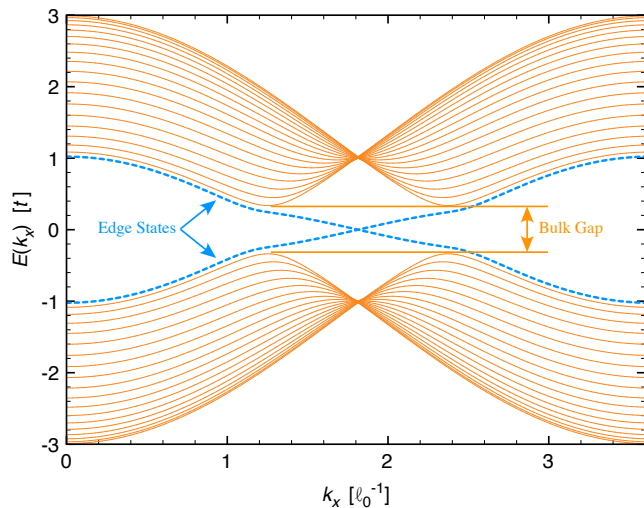


FIG. 3 (Colours online) Edge states of the quantum spin-Hall effect in zig-zag graphene nano ribbons, the bulk states are given by the solid-orange lines, the spin-polarised states are the dashed-blue lines.

Aharonov-Casher (AC) effects (Akkermans and Montambaux, 2007; Altland and D., 2010; Nazarov and Blanter, 2013). This is very interesting because it would allow to realise novel functionalities such as possible non-Abelian operations on spins.

### E. Weyl semimetals

Weyl semimetals also known as “topological semimetals” are zero gap semiconductors with a Fermi level very close to the centre of the gap that is at zero energy as in the case of SLG and TIs. The effective Hamiltonian describing the low energy states close to the Fermi level in WSs is represented by a  $2 \times 2$  Hamiltonian that is linear in the momentum — as for TI and SLG — but it is an actual 3D model, thus containing all the three Pauli matrices. The simplest model Hamiltonian in  $k$   $p$  approximation reads:

$$\mathcal{H}_{\pm} = \pm v_F \mathbf{p} \cdot \mathbf{s}, \quad (7)$$

where  $v_F$  is the Fermi velocity, and  $\mathbf{p}$  and  $\mathbf{s}$  are the vectors of the momentum and of the Pauli matrices, respectively. By inspection of equation 7 we evince that a generic perturbation cannot gap the energy spectrum. Its energy spectrum reads  $E_{\pm}(p) = \pm v_F |\mathbf{p}|$ . This is usually correct for an even number of *nodal* points where the expansion (7) is allowed. At each node we can associate a chirality, this measures the relative handedness of the three momenta and the Pauli matrices associated in the Weyl equation — the chirality  $\pm 1$  for the Hamiltonians  $\mathcal{H}_{\pm}$ , can also be thought as a source of Berry curvature carried by the WS nodes. This is a general property of Weyl fermions realised in band structures: their net chirality

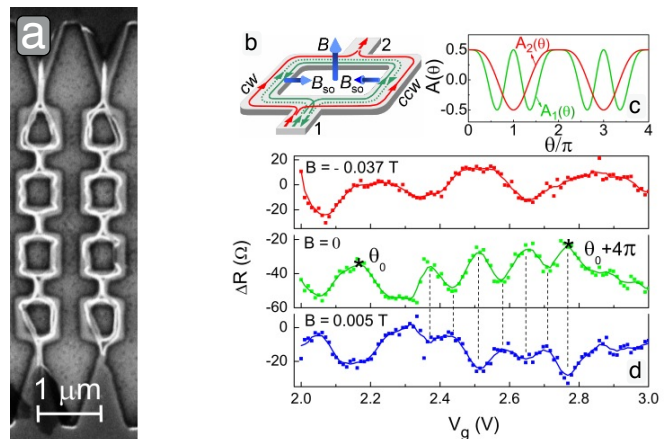


FIG. 4 (Colours online) (a): SEM image of the square-ring device realised in  $\text{Bi}_2\text{Se}_3$ ; (b): Illustration of the AB effect interference (red trajectories) and AAS interference (solid and dotted green loops) for charges in a square ring. The existence of RSOI creates an effective magnetic field  $B_{\text{SO}}$  pointing towards/ or from the center of the ring for counterclockwise (CCW)/clockwise (CW) propagation modes, which influences spin precession and generates an AC phase in addition to the AB and AAS phases. (c)  $A_1(\theta)$  and  $A_2(\theta)$  as a function of  $\theta$ . Here  $\theta = 2\alpha m^* L / \hbar^2$  is spin precession angle over a distance  $L$  and  $A_1(\theta) = (\cos^4 \theta + 4 \cos \theta \sin^2 \theta + \cos 2\theta)/4$  and  $A_2(\theta) = (\sin^2 \theta + \cos 2\theta)/4$  are the modulation of the wave function as a function of  $\theta$  (c.f. Ref. (Qu *et al.*, 2011) for more details). The former varies roughly at twice frequency of the latter. (d):  $\Delta R - V_g$  curves at fixed magnetic fields marked by the lines and arrows of corresponding colours. The spin precession angle is modulated by  $4\pi$  by varying  $V_g$  from the interval 2.16 to 2.77 V marked by the two stars. The dashed lines help illustrating the opposite phases between the green and blue curves in the AAS region. Courtesy of Ref. [Qu *et al.*, 2011].

must in fact cancel<sup>3</sup>. Note that it is usually assumed that bands are individually nondegenerate. This requires that either the time-reversal symmetry or the inversion symmetry (parity) is broken. In order to realise the minimal case of just a pair of opposite chirality Weyl nodes, time-reversal symmetry must be broken (Wan *et al.*, 2011).

The idea of WS was first theoretically proposed by Murakami in 2007 [Murakami, 2007] and later it was further elaborated by Wan *et al.* in 2011 [Wan *et al.*, 2011]. Based on *ab-initio* band calculations, it was proposed that pyrochlore iridates, such as  $\text{Y}_2\text{Ir}_2\text{O}_7$ , in the antiferromagnetic phase, may realise such a WS [Wan *et al.*, 2011 and Yang and Kim, 2010]. A recent review on the spectral properties of this type of system is reported by Vafeek and Vishwanath [Vafeek and Vishwanath, 2014] and on the transport properties by Hosur and Qi [Hosur and Qi, 2013].

<sup>3</sup> This can be understood in terms of the fermion doubling theorem (Nielsen and Ninomiya, 1981).

### III. SPIN TRANSPORT IN RSOI MATERIAL

Before describing the role of RSOI in the spin-dependent transport, it is important to understand how it modifies the spectral properties of a free electron in a 2DEG.

#### A. General properties of the RSOI in semiconductors

We consider a 2DEG in the  $(x, y)$ -plane in the presence of the RSOI (5) and with effective electron mass  $m^*$ . The full Hamiltonian of the system is

$$\mathcal{H}_0 = \frac{\mathbf{p}^2}{2m^*} + \frac{\alpha}{\hbar}(\mathbf{s} \times \mathbf{p})_z. \quad (8)$$

The eigenvalues read

$$\mathcal{E}_{\pm}(\mathbf{k}) = \frac{\hbar^2 k^2}{2m^*} \pm \alpha k = \frac{\hbar^2}{2m^*} (k \pm k_{\text{SO}})^2 - \Delta_{\text{SO}}, \quad (9)$$

where  $k = |\mathbf{k}| = \sqrt{k_x^2 + k_y^2}$  is the modulus of the electron momentum,  $k_{\text{SO}} = \frac{\alpha m^*}{\hbar^2}$  is the RSOI coupling constant with the dimension of a momentum and  $\Delta_{\text{SO}} = \left(\frac{\alpha m^*}{\hbar}\right)^2$ . Usually the last term of (9) is neglected because it is second order in  $\alpha$ . Moreover, even if included, it would lead to a rigid shift of the bands, thus renormalising the chemical potential  $\mu$ . The eigenvectors of the Hamiltonian (8) relative to the spectrum (9) are plane waves:

$$\Psi_{\pm}(\mathbf{r}) = \frac{e^{i\mathbf{k}\cdot\mathbf{r}}}{\sqrt{2}} \begin{pmatrix} 1 \\ \pm i e^{-i\theta} \end{pmatrix}, \quad (10)$$

where  $\theta = \arctan(k_y/k_x)$  is the polar angle between the momentum vector and the  $k_x$  direction. It is important to note that the spin states (10) are always perpendicular to the motion direction. In fact, if an electron moves along  $x$  direction the spinor part of the eigenvectors become  $(1; i)$  and  $(1; -i)$  that is the spin up and spin down are along the  $y$  direction. By contrast, if the electron moves along the  $y$  direction, the eigenvectors become  $(1; 1)$  and  $(1; -1)$  that is the spin up and spin down state in the  $x$  direction [see figure 5(b)].

In figure 5(c)-(e) we report the  $k_y$ -section of the energy spectrum *vs* the momentum for a 2DEG in different physical situations. In 5(c) we considered a free particle in a 2DEG. In this case the spectrum is twofold degenerate in spin. In the presence of a magnetic field  $\mathbf{B}$  [see panel 5(d)], the spin degeneracy is lifted out by the *Zeeman* effect and the gap separating spin up and spin down is equal to  $2g^*\mu_B B$ , where  $g^*$  is the effective gyromagnetic ratio and  $\mu_B$  is the Bohr's magneton. When the RSOI is present 5(e), the spin degeneracy is lifted out except for  $k_y = 0$ . In this situation the degeneracy is removed without the opening of gaps. The semiclassical particle

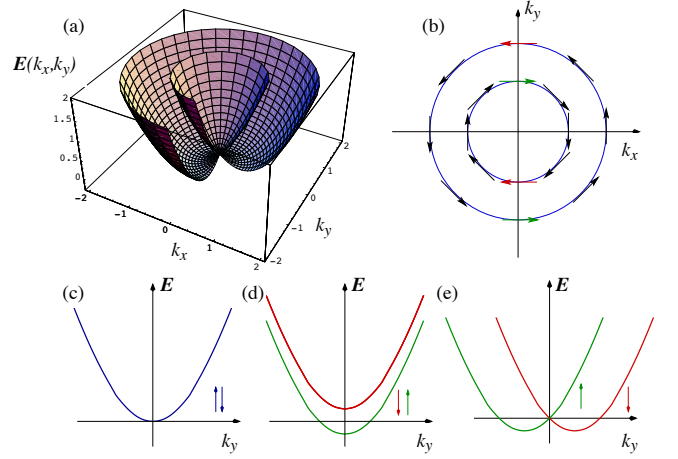


FIG. 5 (Colours online) Properties of the Rashba energy spectrum. (a) Portion of the energy spectrum of the Hamiltonian (8). (b) The Fermi contours relative to the Hamiltonian (8), the spin states are shown as well. (c) Section of the energy spectrum for a free electron. (d) Section of the energy spectrum for an electron in presence of a magnetic field, *e.g.* *Zeeman* splitting. (e) Section of the energy spectrum for an electron in presence of RSOI.

velocities are given by

$$\mathbf{v}_{\pm}(\mathbf{k}) = \frac{1}{\hbar} \frac{\partial \mathcal{E}_{\pm}(\mathbf{k})}{\partial \mathbf{k}} = \frac{\hbar \mathbf{k}}{m^*} \pm \alpha \mathbf{k} = \frac{\hbar}{m^*} (k \pm k_{\text{SO}}) \mathbf{k}. \quad (11)$$

If we consider the quantum-mechanical velocity operator

$$\hat{\mathbf{r}} = \frac{i}{\hbar} [\mathcal{H}_0, \mathbf{r}] \quad (12)$$

and the expression (10) for the eigenstates, it is straightforward to show that its matrix elements are given by

$$\langle \Psi_{\pm}(\mathbf{k}) | \hat{\mathbf{r}} | \Psi_{\pm}(\mathbf{k}') \rangle = \delta(\mathbf{k} - \mathbf{k}') \mathbf{v}_{\pm}(\mathbf{k}). \quad (13)$$

That is the semiclassical velocities  $\mathbf{v}_{\pm}(\mathbf{k})$  are, as usual, the diagonal elements of the velocity operator. In the presence of the RSOI, the velocity operator is not simply the momentum divided by the effective mass as for free electrons and the Fermi surface splits into two surfaces shown in figure 5(b). One can easily observe that counter propagating electrons have opposite spins. Spin and momentum are locked to each other. This will have importance consequences in the following. Parametrising the wave vectors as  $\mathbf{k} = k (\cos \varphi, \sin \varphi)$  the two Fermi surfaces are described by the following equation:

$$k_{\pm}^{\text{F}}(\mathcal{E}_{\text{F}}) = \mp \frac{\alpha m^*}{\hbar^2} + \sqrt{\left(\frac{\alpha m^*}{\hbar^2}\right)^2 + \frac{2m^*}{\hbar^2} \mathcal{E}_{\text{F}}}. \quad (14)$$

Here the double sign corresponds to the two dispersion branches (9).

## B. Rashba SOI in quantum wires: subbands hybridisation

By further confining a 2DEG along one direction (we choose  $y$  in the following) we can realise a quantum wire. In this case we cannot solve analytically the system Hamiltonian, as we have done for the simple case of a 2DEG. Different theoretical models have been used to describe confinement effects. A very convenient representation consists in assuming a transversal confining potential in the  $y$  directions, and let the electrons propagate along the  $x$  direction. Such choice is particularly convenient when expressing the matrix elements of the SOI (Moroz and Barnes, 1999; Governale and Zülicke, 2002; Perroni *et al.*, 2007). The Hamiltonian reads:

$$H = \mathcal{H}_{\parallel} + \mathcal{H}_{\perp} + \mathcal{H}_{\text{mix}}, \quad (15)$$

with the following terms:

$$\mathcal{H}_{\parallel} = \frac{p_x^2}{2m^*} - \frac{\hbar k_{\text{SO}}}{m^*} \sigma_y p_x \quad (16a)$$

$$\mathcal{H}_{\perp} = \frac{p_y^2}{2m^*} + V(y) \quad (16b)$$

$$\mathcal{H}_{\text{mix}} = \frac{\alpha}{\hbar} \sigma_x p_y. \quad (16c)$$

Where  $V(y)$  is a infinite well potential of width  $W$ . Neglecting  $\mathcal{H}_{\text{mix}}$ , the terms in equations (16a) and (16b) do commute, therefore the eigenvalues and eigenvectors can be easily expressed as:

$$E_{\parallel+\pm}(n, \sigma_y, k_x) = \frac{\hbar^2}{2m^*} \left( k_x^2 + \frac{\pi^2 n^2}{W^2} \right) - \frac{\hbar^2 k_{\text{SO}} k_x}{m^*} \langle \sigma_y \rangle \quad (17)$$

$$\psi_{n\sigma_y k_x}(x, y) = \phi_n(y) \psi_{k_x}(x) |\sigma_y\rangle \quad (18)$$

where  $\phi_n(y)$  are the  $n$ -th eigenfunctions of the potential  $V(y)$  that are either a sine or a cosine functions and  $\psi_{k_x}(x)$  simple plane waves shifted by the RSOI, here  $\langle \sigma_y \rangle$  can get the values  $\pm 1$ . The term  $\mathcal{H}_{\text{mix}}$  induces mixing between these states and gives rise to a deformation of the electronic bands and to anti-crossings in the energy spectrum.

Anti-crossings occur between sub-bands corresponding to transverse eigenstates with different  $n$  [figure 6(a)]. While simple one dimensional (1D) models predict a rigid spin-momentum locking with  $in$ -plane spin perpendicular to the momentum along the wire direction, multi-band models predict that only electrons with momentum far away from anti-crossings have the spin essentially perpendicular to the momentum, *i.e.*  $\langle S_y \rangle \sim \pm \frac{1}{2}$  and  $\mathbf{k} \sim k_x \mathbf{x}$ . This is well shown in figure 6(b), where we can see the expectation value of  $\langle S_y \rangle$  as a function of the longitudinal momentum  $k_x$  — it changes from  $-\frac{1}{2}$  to  $\frac{1}{2}$  for  $k_x$  changing from negative to positive values. The same picture applies to the other sub-bands. Here we shall focus on the role played by the number of sub-bands used for evaluating the system properties. The

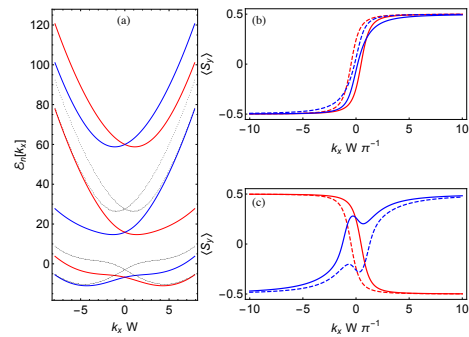


FIG. 6 (Colours online) (a) Band structure of quantum wires with square well potential along the confining direction. The coloured lines refer to the first three sub-bands in the case of a wire with three modes whereas the dotted-black lines refer to the same wire but only considering two modes. (b) Spin polarisation  $\langle S_y \rangle$  as a function of the transverse momentum for the first sub-band in the case of a two band model (red-solid and red-dashed lines). (c) Spin polarisation  $\langle S_y \rangle$  as a function of the transverse momentum for the second sub-band in the case of a two band model (red-solid and red-dashed lines). Both in panel (b) and (c) the spin polarisation is evaluated using a truncated Hilbert space containing 50 modes (blue-solid and blue-dashed lines).

Hamiltonian (15) cannot be diagonalized exactly, but a partial analytical/numerical solution can be obtained by truncating the Hilbert space associated to the modes produced by  $V(y)$ . Diagonalizing the Hamiltonian (15) for  $N \gg 1$  modes produces a result that is substantially correct for the lowest  $N - 1$  modes but not for the mode  $N$ . This is observed, *e.g.*, in the spectral properties and in the spin polarisation. The spectrum is shown in figure 6(a), we have in colours-solid lines for  $N = 3$  and in black-dashed lines for  $N = 2$ . We see that the first sub-bands in both cases ( $N = 3$  and  $N = 2$ ) are quite similar apart from a negative energy shift (Perroni *et al.*, 2007). However, the situation is very different when we consider the second sub-bands, these greatly differ in the two cases — due to the presence of the third mode for the case  $N = 3$ . Considering the spin polarisation the effect is more visible. In figure 6(b) the spin polarisations for the first sub-bands for  $N = 2$  (red lines) and  $N = 50$  (blue lines) are shown, the behaviours are similar in both cases but around  $k_x \sim 0$ . The error produced by Hilbert space truncation is shown in 6(c), here it is shown the spin polarisation for the second sub-bands (red lines), for  $N = 2$ . We observe a mirroring of the results reported in 6(b). However, the most correct result with  $N = 50$  (blue lines) contradicts completely  $N = 2$ .

The message here is that when solving a problem of quantum transport in a confined geometry a sufficient number of sub-bands should be taken into account in order to avoid systematic errors due to the truncation of the Hilbert space (Moroz and Barnes, 1999; Governale and Zülicke, 2002; Perroni *et al.*, 2007). The same con-



clusions are obtained by changing the shape of the confining potential  $V(y)$  from square potential one to harmonic oscillator one (Moroz and Barnes, 1999; Governale and Zülicke, 2002) or by introducing a further periodic modulation along the longitudinal direction  $x$  (Smirnov *et al.*, 2007). The sub-band hybridisation that we have shown before can give rise to dip into the conductance of quantum wires with RSOI as shown in several works both analytically (Moroz and Barnes, 1999; Perroni *et al.*, 2007) and numerically (Sánchez and Serra, 2006) and can also be interpreted in terms of a Landau-Zener transition when analysing the conductance of a quantum point contact (Eto Mikio *et al.*, 2005).

### C. General properties of the Rashba SOI in graphene

If we restrict ourselves to the case of a single valley approximation, the effects of RSOI on SLG via Hamiltonian (6) change with respect to what we have seen in the case of a 2DEG in Sec. III.A. The complete Hamiltonian for the SLG with RSOI reads:

$$\begin{aligned} \mathcal{H}_{\mathbf{K}} &= v_{\text{F}} \boldsymbol{\sigma} \cdot \mathbf{p} + \frac{\alpha}{2} (\mathbf{s} \times \boldsymbol{\sigma})_z \\ &= v_{\text{F}} (\sigma_x p_x + \sigma_y p_y) + \frac{\alpha}{2} (s_x \sigma_y - s_x \sigma_y). \end{aligned} \quad (19)$$

The energy eigenstates are plane waves  $\psi \sim \Phi(\mathbf{k})e^{i\mathbf{k}\cdot\mathbf{r}}$  with  $\Phi$  a four-component spinor and eigenvalues given by ( $v_{\text{F}} = \hbar = 1$ )

$$\mathcal{E}_{\pm, \epsilon}(\mathbf{k}) = \pm \frac{\alpha}{2} + \epsilon \sqrt{k_x^2 + k_y^2 + \frac{\alpha^2}{4}}, \quad (20)$$

where index  $\epsilon = \pm$  specifies the particle/hole branches of the spectrum. The energy dispersion, as a function of  $k_x$ , at fixed  $k_y = 0$ , is illustrated in figure 7 with and without RSOI. As we can see here, the RSOI gives rise to a finite

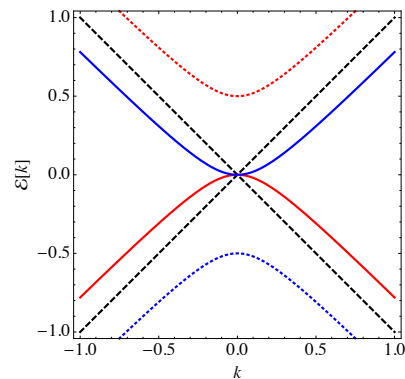


FIG. 7 (Colours online) Energy spectrum of graphene for  $k_y = 0$  as a function of  $k$  along the  $x$  axis. The dashed lines correspond to the linear dispersion for  $\alpha = 0$ , the solid and the dotted lines to the case of finite RSOI. The lines with the same colours correspond to the same spin state.

curvature of the linear energy spectrum and lifts the spin degeneracy. Two opposite spin bands get gapped while the other two are still degenerate at  $\mathbf{k} \sim 0$ . However, contrary to the case of 2DEGs, the energy spectrum is not shifted along the momentum axis as in equation (9). This is related to the fact that the RSOI Hamiltonian does not depend on the electron momentum but only on its pseudo-spin, to the lowest order in the momentum expansion. Nevertheless, this approximation no longer holds if we expand RSOI Hamiltonian to higher orders, as they explicitly depend on the electron momentum. Their main effect, for strong RSOI, is to produce the appearing of extra Dirac cones (tridiagonal wrapping Ref. [Zarea and Sandler, 2009; Rakyta *et al.*, 2010; and Lenz *et al.*, 2013]).

The eigenstates of the Hamiltonian (19), are expressed by the spinors  $\Phi_{\pm, \epsilon}(\mathbf{k})$ :

$$\Phi_{\pm, \epsilon}^{\text{T}}(\mathbf{k}) = \frac{1}{2\sqrt{\cosh \theta_{\pm}}} \times (e^{-i\phi - \epsilon\theta_{\pm}/2}, \epsilon e^{\epsilon\theta_{\pm}/2}, \pm i\epsilon e^{\epsilon\theta_{\pm}/2}, \pm i e^{i\phi - \epsilon\theta_{\pm}/2}),$$

where  $\square^{\text{T}}$  denotes transposition and

$$\sinh \theta_{\pm} = \pm \frac{\alpha}{2k}, \quad (21)$$

$$e^{i\phi} = \frac{k_x + ik_y}{k}, \quad (22)$$

with  $k = \sqrt{k_x^2 + k_y^2}$ . The spin operator components are expressed as  $S_j = \frac{1}{2} s_j \otimes \mathbb{I}_2$ . Their expectation values

on the eigenstates  $\Phi_{\pm, \epsilon}$  read

$$\langle S_x \rangle = -\frac{\epsilon \sin \phi}{2 \cosh \theta_{\pm}}, \quad (23a)$$

$$\langle S_y \rangle = \frac{\epsilon \cos \phi}{2 \cosh \theta_{\pm}}, \quad (23b)$$

$$\langle S_z \rangle = 0, \quad (23c)$$

which shows that the product  $\pm\epsilon$  coincides with the sign of the expectation value of the spin projection along the  $in$ -plane direction perpendicular to the direction of propagation. For vanishing RSOI, the eigenstates  $\Phi_{\pm, \epsilon}$  reduce to linear combinations of eigenstates of  $S_z$ . Similarly, the

expectation value of the pseudo-spin operator  $\boldsymbol{\sigma}$  is given by

$$\langle \sigma_x \rangle = \frac{\epsilon \cos \phi}{\cosh \theta_{\pm}}, \quad (24a)$$

$$\langle \sigma_y \rangle = \frac{\epsilon \sin \phi}{\cosh \theta_{\pm}}. \quad (24b)$$

Since the RSOI in SLG does not depend on momentum, the velocity operator still coincides with the pseudo-spin operator:  $\mathbf{v} = \dot{\mathbf{r}} = i[\mathcal{H}, \mathbf{r}] = \boldsymbol{\sigma}$ . Thus, the velocity expectation value in the state  $\Phi_{\pm, \epsilon}$  is given by equations (24a) and (24b). Alternatively, it can be obtained from the energy dispersion as

$$\mathbf{v}_{\epsilon} = \nabla_{\mathbf{k}} \mathcal{E}_{\epsilon} = \frac{\epsilon \mathbf{k}}{\sqrt{k^2 + \frac{\alpha^2}{4}}}. \quad (25)$$

If we considered also the intrinsic SOI, the effective Fermi velocity would acquire a more complex dependence from the RSOI (Lenz *et al.*, 2013; Bercieux and De Martino, 2010).

Sub-band hybridisation due to RSOI also occurs in graphene nano-ribbons. However, its manifestation depends on the type of boundary (Neto *et al.*, 2009): zig-zag (Zarea and Sandler, 2009) or armchair (Lenz *et al.*, 2013). The most peculiar effects are observed in the case latter. Here in fact, the RSOI couples not only different modes but introduces also a finite coupling inside each sub-band. It is due to the presence of two inequivalent carbon atoms in the unit cell of a SLG nano ribbon and is completely different from the case of a quantum wire with RSOI of the previous section (Lenz *et al.*, 2013).

#### D. The spin double refraction

The physics of an interface between a 2DEG with and without SOI is very similar to that of the optical birefringence (Born and Wolf, 1999) and is a direct consequence of the two possible Fermi velocities for the two modes (11). In the following we consider a 2DEG in the  $(x, y)$ -plane characterised by an effective electron mass  $m^*$  and an interface along the  $y$  direction separating a region without SOI (N-region) and a region with it (SO region). Elastic scattering at the interface allows for conservation of the total energy and of the momentum parallel to the interface —  $k_y$  in our case. As we have seen in the previous section, the Fermi surface for an energy  $\mathcal{E}_F$  is constituted of two circles with radius (14). An incoming particle from the N region is characterised by the momentum  $k^N$  and the incidence angle  $\phi = \arctan\left(\frac{k_y^N}{k_x^N}\right)$ . Momentum conservation implies that

$$k_+^{\text{SO}} \sin \theta_+ = k_-^{\text{SO}} \sin \theta_- = k^N \sin \phi, \quad (26)$$

where we have introduced  $k_{\pm}^{\text{SO}} = k_{\pm}^F(\mathcal{E}_F)$  and  $\theta_{\pm}$  are the propagation angles in the SO region for the two different

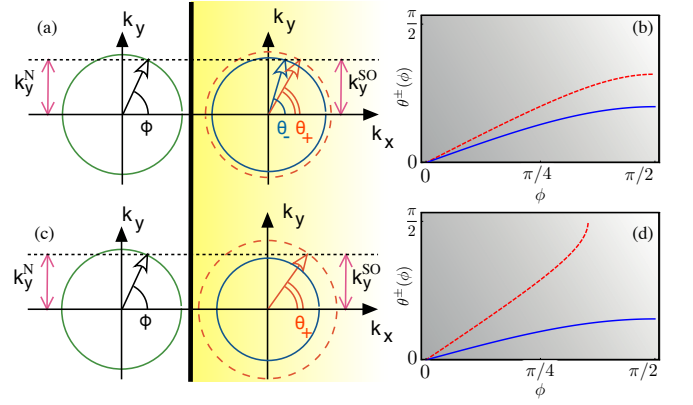


FIG. 8 (Colours online) Spin Refraction angles  $\theta_{\pm}$  as a function of the incoming particle angle  $\phi$ . Panel (a) and (b) case parameters of two modes always open. Panel (c) and (d) case parameters where the critical angle of mode  $(-)$ ,  $\phi_c^-$  is bigger than  $\pi/2$ . In Panels (b) and (c) the red-dashed line is the mode  $(-)$  whereas the blue-solid line is the mode  $(+)$ .

modes. An incoming particle from the normal region propagates into the SO region along two different modes with indices  $\pm$ . In Refs. [Perroni *et al.*, 2007; Ramaglia *et al.*, 2003; Khodas *et al.*, 2004; and Ramaglia *et al.*, 2004] this phenomenon is named *spin-double refraction*. From equation (26) we can obtain an expression for the two refraction angles:

$$\theta_{\pm} = \arcsin\left(\frac{k^N}{k_{\pm}^{\text{SO}}} \sin \phi\right). \quad (27)$$

If the refraction angle is equal to  $\frac{\pi}{2}$ , then the corresponding mode will not propagate forward, thus it is *closed*. Of course, there are two critical angles  $\phi_c^{\pm}$  corresponding to the closure of the corresponding modes in the SO region defined by the relation:

$$\phi_c^{\pm} = \arcsin\left(\frac{k_{\pm}^{\text{SO}}}{k^N}\right). \quad (28)$$

According to equation (14):  $k_+^{\text{SO}} < k_-^{\text{SO}}$  so that  $\phi_c^+ < \phi_c^-$ . Furthermore, considering that  $k_+^{\text{SO}}$  is always smaller than  $k^N$ , the mode  $(+)$  will be always open. The previous results are based only on kinematic considerations and are independent by the nature of the SOI. The key ingredient is the presence of an interaction that splits the 2D Fermi surface in two circles. For example, the same physics would be possible at an interface between a normal region and a region with a strong Zeeman splitting, assuming that the Fermi energy allows the propagation along the two modes in the region with the magnetic field.

Some care is required when writing down the Hamiltonian for this scattering problem. Indeed the two regions (N and SO) are obtained by considering an inhomogeneous coupling constant for the RSOI  $\alpha(x)$ . As a consequence,  $\alpha(x)$  does not commute with the momentum

operator  $[\alpha(x), p_x] \neq 0$ . Thus the Hamiltonian describing the two regions has to read:

$$\mathcal{H}_{\text{N-SO}} = \frac{\mathbf{p}^2}{2m^*} + \frac{\alpha(x)}{\hbar}(s_x p_y - s_y p_x) - i s_y \frac{1}{2m^*} \frac{\partial \alpha(x)}{\partial x} + V_{\text{int}} \delta(x). \quad (29)$$

where the second to last terms reestablish the hermiticity (Messiah, 2014). In the simplest approximation we can consider  $\alpha(x) = \alpha \Theta(x - x_{\text{int}})$ , where  $\Theta(x)$  is the Heaviside step function. The last term mimics the presence an SOI interface, according to the standard approach describing interfaces with delta potentials of strength  $V_{\text{int}}$  (Blonder *et al.*, 1982). A more realistic description has to account also for possible changes of the effective electron mass  $m^* \rightarrow m^*(x) = m_{\text{N}}^* \Theta(x_{\text{int}} - x) + m_{\text{SO}}^* \Theta(x_{\text{int}} + x)$  in the two regions, more details can be found in Ref. [Perroni *et al.*, 2007 and Ramaglia *et al.*, 2003]. When considering the presence of a second interface, where the RSOI vanishes again, the refraction angles for the two modes will be the same and equal to the incoming one  $\phi$  (Khodas *et al.*, 2004; Ramaglia *et al.*, 2004). However this double interface structure cannot produce a spin polarisation when the incoming particles are unpolarised. This is due to the fact that in order to produce a spin polarisation we need to violate the Onsager relation for a two terminal system (Adagideli *et al.*, 2012; Gorini *et al.*, 2012). Violation is possible if and only if time reversal symmetry is broken, this implies, *e.g.*, the application of a magnetic field.

The same physics is possible also in the case of SLG, the main difference is that the RSOI in this case does not contain the momentum operator (6). As a consequence we do not need the extra term in the full Hamiltonian as in equation (29). A complete analysis of the spin-double refraction for the case of graphene is reported in Ref. (Bercioux and De Martino, 2010) where a transfer matrix method is presented, which is useful for studying the spin-dependent transport in hybrid structures in SLG, as a periodic RSOI potential (Lenz and Bercioux, 2011; Costa *et al.*, 2013).

#### IV. PURE SPIN CURRENT GENERATION IN RSOI MATERIAL

A very crucial point in spintronics is the creation of pure spin currents. Over the last 20 years, many methods for creating pure spin currents have been proposed. We will focus mainly on ratchet and pumping methods that have received also experimental verification. Here we define what is a spin current and discuss a method for evaluating it in terms of the Landauer-Büttiker formalisms (Scheid *et al.*, 2007a). A pure spin current is defined as a particle flow carrying finite spin polarisation without an associated charge current.

A simple example follows: Suppose, for instance, that a charge  $\mathcal{Q}$  and spin polarisation  $\mathcal{S}_{pq}$  moves from a contact  $p$  to  $q$  during the time  $t \in [0, T]$  and in the next period  $t \in [T, 2T]$  the same amount of charge  $\mathcal{Q}$ , but a different spin polarisation  $\mathcal{S}_{qp}$ , moves from contact  $q$  to  $p$ . The net charge transported between  $p$  and  $q$ , in the time  $t \in [0, 2T]$  is zero, but the net spin polarisation is  $\Delta \mathcal{S} = \mathcal{S}_{pq} - \mathcal{S}_{qp}$ .

In order to evaluate the spin-current we consider  $N$  non ferromagnetic contacts injecting spin-unpolarised current into the leads. We use, as customary, a local coordinate system for the lead under investigation, where  $x$  is the coordinate along the lead in the direction of charge propagation due to an applied bias in linear response and  $y$  is the transverse coordinate. In each lead, at a fixed energy  $E$ , several conducting modes are open. The wave function for each mode reads

$$\Psi_{E,ns}^{\pm}(x, y) = \frac{1}{\sqrt{k_n(E)}} e^{\pm i k_n(E)x} \chi_n(y) \Sigma(s), \quad (30)$$

here we have introduced the transverse eigenfunctions of the lead of width  $W$ :

$$\chi_n(y) = \sqrt{\frac{2}{W}} \sin\left(\frac{n\pi y}{W}\right) \quad \forall n \in \mathbb{N}$$

with the eigenenergy  $E_n = \frac{\hbar^2}{2m^*} \left(\frac{n\pi}{W}\right)^2$  and  $\Sigma(s)$  is the spin eigenfunction. The superscript  $\pm$  of  $\Psi$  refers to the motion direction along the lead axis with the wave-vector  $k_n = \sqrt{2m^*(E - E_n)}/\hbar$ . We use the scattering approach for deriving the current formula: the amplitudes of the states inside the leads are related via the scattering matrix  $\mathcal{S}(E)$ , determined by the Hamiltonian of the coherent conductor. The scattering state inside the  $q$ -th lead reads

$$\varphi_E^q(x, y) = \sum_{ns} \left[ a_{ns}^q(E) \Psi_{E,ns}^+(x, y) + b_{ns}^q(E) \Psi_{E,ns}^-(x, y) \right],$$

with ( $s = \pm$ ). It consists of incoming states  $\Psi^+$  entering the coherent conductor from the contact  $q$  and outgoing states  $\Psi^-$  leaving the coherent conductor into the contact  $q$ . The amplitudes of incoming  $a_{ns}^j$  and outgoing  $b_{ns}^i$  states are related each other via the scattering relation

$$b_{n's'}^i(E) = \sum_{j=1}^N \sum_{n \in j} \sum_{s=\pm 1} S_{n's',ns}^{i,j}(E) a_{ns}^j(E), \quad (31)$$

where the scattering matrix  $\mathcal{S}(E)$  has the following structure for an  $N$  terminal system:

$$\mathcal{S}(E) = \begin{pmatrix} \mathbf{r}^{1,1}(E) & \mathbf{t}^{1,2}(E) & \dots & \mathbf{t}^{1,N}(E) \\ \mathbf{t}^{2,1}(E) & \mathbf{r}^{2,2}(E) & \dots & \mathbf{t}^{2,N}(E) \\ \vdots & \vdots & \ddots & \vdots \\ \mathbf{t}^{N,1}(E) & \mathbf{t}^{N,2}(E) & \dots & \mathbf{r}^{N,N}(E) \end{pmatrix}.$$

Here the sub-matrix  $\mathbf{r}^{j,j}(E)$  is a square matrix of dimensionality  $M^j(E)$ , corresponding to the number of open channels in lead  $j$  at energy  $E$  connected to a reservoir with chemical potential  $\mu_j$  — it already includes the spin degree of freedom. The matrix  $\mathbf{r}^{j,j}(E)$  contains the scattering amplitudes of incoming channels of lead  $j$  being reflected back into outgoing channels of the same lead. The sub-matrix  $\mathbf{t}^{i,j}(E)$  is a  $M^i(E) \times M^j(E)$  matrix that contains the scattering amplitudes for transmission between incoming channels from lead  $j$  and outgoing channels of lead  $i$ .

In order to evaluate the spin-current we consider  $N$  non ferromagnetic contacts injecting spin-unpolarised current into the leads. We use, as customary, a local coordinate system for the lead under investigation, where  $x$  is the coordinate along the lead in the direction of charge propagation due to an applied bias in linear response and  $y$  is the transverse coordinate. In each lead, at a fixed energy  $E$ , several conducting modes are open. The wave function for each mode reads

$$\Psi_{E,ns}^{\pm}(x,y) = \frac{1}{\sqrt{k_n(E)}} e^{\pm i k_n(E)x} \chi_n(y) \Sigma(s), \quad (32)$$

here we have introduced the transverse eigenfunctions of the lead of width  $W$ :

$$\chi_n(y) = \sqrt{\frac{2}{W}} \sin\left(\frac{n\pi y}{W}\right) \quad \forall n \in \mathbb{N}$$

with the eigenenergy  $E_n = \frac{\hbar^2}{2m^*} \left(\frac{n\pi}{W}\right)^2$  and  $\Sigma(s)$  is the spin eigenfunction. The superscript  $\pm$  of  $\Psi$  refers to the motion direction along the lead axis with the wave-vector  $k_n = \sqrt{\frac{2m^*}{\hbar^2}(E - E_n)}$ . We use the scattering approach for deriving the current formula: the amplitudes of the states inside the leads are related via the scattering matrix  $\mathbf{S}(E)$ , determined by the Hamiltonian of the coherent conductor. The scattering state inside the  $q$ -th lead reads

$$\varphi_E^q(x,y) = \sum_{ns} \left[ a_{ns}^q(E) \Psi_{E,ns}^+(x,y) + b_{ns}^q(E) \Psi_{E,ns}^-(x,y) \right],$$

with ( $s = \pm$ ). It consists of incoming states  $\Psi^+$  entering the coherent conductor from the contact  $q$  and outgoing states  $\Psi^-$  leaving the coherent conductor into the contact  $q$ . The amplitudes of incoming  $a_{ns}^j$  and outgoing  $b_{ns}^i$  states are related each other via the scattering relation

$$b_{n's'}^i(E) = \sum_{j=1}^N \sum_{n \in j} \sum_{s=\pm 1} S_{n's',ns}^{i,j}(E) a_{ns}^j(E), \quad (33)$$

where the scattering matrix  $\mathbf{S}(E)$  has the following structure for an  $N$  terminal system:

$$\mathbf{S}(E) = \begin{pmatrix} \mathbf{r}^{1,1}(E) & \mathbf{t}^{1,2}(E) & \dots & \mathbf{t}^{1,N}(E) \\ \mathbf{t}^{2,1}(E) & \mathbf{r}^{2,2}(E) & \dots & \mathbf{t}^{2,N}(E) \\ \vdots & \vdots & \ddots & \vdots \\ \mathbf{t}^{N,1}(E) & \mathbf{t}^{N,2}(E) & \dots & \mathbf{r}^{N,N}(E) \end{pmatrix}.$$

Here the sub-matrix  $\mathbf{r}^{j,j}(E)$  is a square matrix of dimensionality  $M^j(E)$ , corresponding to the number of open channels in lead  $j$  at energy  $E$  connected to a reservoir with chemical potential  $\mu_j$  — it already includes the spin degree of freedom. The matrix  $\mathbf{r}^{j,j}(E)$  contains the scattering amplitudes of incoming channels of lead  $j$  being reflected back into outgoing channels of the same lead. The sub-matrix  $\mathbf{t}^{i,j}(E)$  is a  $M^i(E) \times M^j(E)$  matrix that contains the scattering amplitudes for transmission between incoming channels from lead  $j$  and outgoing channels of lead  $i$ .

The wave function of the scattering state inside lead  $i$ , where only the incoming channel ( $ns$ )  $\in j$  is populated ( $a_{n's'}^j = \delta_{j',j} \delta_{n',n} \delta_{s',s}$ ), reads for  $j = i$ :

$$\varphi_{E,ns}^i(x,y) = \Psi_{E,ns}^+(x,y) + \sum_{(n's') \in i} r_{n's',ns}^{i,i}(E) \Psi_{E,n's'}^-(x,y), \quad (34)$$

and, correspondingly, for  $j \neq i$

$$\varphi_{E,ns}^i(x,y) = \sum_{(n's') \in i} t_{n's',ns}^{i,j}(E) \Psi_{E,n's'}^-(x,y). \quad (35)$$

For a generic spin wave function  $\Phi(x,y)$  the spin current  $I_S(x)$  passing a cross section ( $x = \text{const}$ ) of a lead is given by:

$$I_S(x) = \int dy \Phi^*(x,y) \mathbf{J}_S \Phi(x,y). \quad (36)$$

Here, we have used the most common definition of the spin current operator (Rashba, 2006). This is defined with respect to an arbitrary quantisation axis  $\mathbf{u}$ , than the operator  $\mathbf{J}_S$  reads:

$$\mathbf{J}_S = \left(\frac{\hbar}{2} \mathbf{s} \cdot \mathbf{u}\right) \left[ \frac{\hbar}{2m^* i} \left( \frac{\partial}{\partial x} \Big|_{\rightarrow} - \frac{\partial}{\partial x} \Big|_{\leftarrow} \right) \right]. \quad (37)$$

In this operator we have the spin operator projected along the  $\mathbf{u}$  quantisation direction times that standard quantum mechanical current. Here, as customary, the partial derivatives act on the expressions to their right and left (as indicated by the arrows), respectively. By acting this the former operator on the scattering state (34) we obtain for the spin current (36) inside lead  $i_{n\sigma}$

$$I_{S;E,ns}^{j=i}(x \in i) = \frac{\hbar^2}{2m^*} \left[ s - \sum_{(n's') \in i} s' \left| r_{n's',ns}^{i,i}(E) \right|^2 \right],$$

where ( $n, s \in j, j = i$ ). For the other scattering state (35) we find the corresponding expression ( $n, s \in j, j \neq i$ )

$$I_{S;E,ns}^{j \neq i}(x \in i) = -\frac{\hbar^2}{2m^*} \sum_{(n's') \in i} s' \left| t_{n's',ns}^{i,j}(E) \right|^2.$$

Each channel is populated according to the Fermi-Dirac distribution  $f(E; \mu_n)$  of the respective contact  $n$ , the statistical average of the total spin current in lead  $i$  reads

$$\begin{aligned} \langle I_S(x \in i) \rangle &= \frac{m^*}{2\pi\hbar^2} \int_0^\infty dE \\ &\left[ \sum_{j=1}^N \sum_{(ns) \in j} f(E; \mu_j) I_{S;E,ns}^j(x \in i) \right] \quad (38) \\ &= -\frac{1}{4\pi} \int_0^\infty dE \left[ f(E; \mu_i) R_S^{i,i}(E) \right. \\ &\quad \left. + \sum_{q \neq i} f(E; \mu_q) T_S^{i,q}(E) \right] \end{aligned}$$

where the spin resolved transmission and reflection are defined as

$$\begin{aligned} T_S^{i,q}(E) &= \sum_{s'=\pm} \left( T_{+,s'}^{i,q} - T_{-,s'}^{i,q} \right) \\ R_S^{i,i}(E) &= \sum_{s'=\pm} \left( R_{+,s'}^{i,i} - R_{-,s'}^{i,i} \right), \end{aligned}$$

with

$$T_{s,s'}^{i,q}(E) = \sum_{n \in i} \sum_{n' \in q} \left| t_{ns,n's'}^{i,q}(E) \right|^2, \quad (39)$$

$$R_{s,s'}^{i,i}(E) = \sum_{n \in i} \sum_{n' \in i} \left| r_{ns,n's'}^{i,i}(E) \right|^2. \quad (40)$$

Probability conservation implies that the scattering matrix has to be unitary  $[\mathcal{S}(E)]^\dagger \mathcal{S}(E) = \mathcal{S}(E)[\mathcal{S}(E)]^\dagger = \mathbb{I}$ , than the following relation holds:

$$\sum_{(n's') \in i} \left| r_{ns,n's'}^{i,i}(E) \right|^2 + \sum_{q \neq i} \sum_{(n''s'') \in q} \left| t_{ns,n''s''}^{i,q}(E) \right|^2 = 1.$$

By using the symmetry relation between the spin transmission and reflection (c.f. Ref. [Scheid *et al.*, 2007a]), it is straightforward to show that

$$R_S^{i,i}(E) + \sum_{q \neq i} T_S^{i,q}(E) = 0.$$

In view of Eq. (38) we eventually find for the spin current in lead  $i$

$$I_S(x \in i) = \frac{1}{4\pi} \int_0^\infty dE \sum_{q \neq i} [f(E; \mu_i) - f(E; \mu_q)] T_S^{i,q}(E). \quad (41)$$

Equilibrium spin currents can locally exist in systems with SOI as shown for 2DEGs (Rashba, 2003) and in mesoscopic systems (Nikolić *et al.*, 2006), however equation (41) clearly shows that at thermal equilibrium ( $\mu_j = \mu \quad \forall j \in \{1, N\}$ ) the spin current vanishes inside leads without SOI and/or magnetic field. This absence of equilibrium spin currents in the leads has been shown for systems with preserved time-reversal symmetry (Kiselev and

Kim, 2005). An expression very similar to (41) has been used to investigate the SHE (Hankiewicz *et al.*, 2004; Ren *et al.*, 2006; Bardarson *et al.*, 2007; Brüne *et al.*, 2012). In the presence of SOI or magnetic fields it is nontrivial to write down a continuity equation for the spin current, the problem has been addressed in several articles (Nikolić *et al.*, 2006; Sun and Xie, 2005; Shi *et al.*, 2006; An *et al.*, 2012).

## A. Pure spin current generators

In this section we describe two mechanisms that can be used for generating a pure spin current.

### 1. Spin Ratchet

A particle ratchet (Reimann, 2002; Hänggi and Marchesoni, 2009) is a system with broken inversion (left/right) symmetry that generates a net currents upon external *ac*-driving in the absence of a net (time-averaged) bias potential. Ratchets have much in common with current rectifiers, though there are differences, in particular in the dissipative case (Reimann, 2002). The theoretical concept of ratchets, originally introduced for classical dynamics, was later extended to the quantum dissipative regime (Reimann *et al.*, 1997). The main difference between ratchets and rectifiers is that quantum ratchets exhibit current reversal upon changing, *e.g.*, the temperature or energy (Hänggi and Marchesoni, 2009; Reimann *et al.*, 1997). Such quantum ratchets were experimentally realised in semiconductor heterostructures in a chain of asymmetric ballistic electron cavities in the low-temperature regime, with a dynamics was close to coherent (Linke *et al.*, 1999). Also the charge current reversal phenomenon has been demonstrated (Linke *et al.*, 2002).

A spin ratchet is a quantum ratchet with *lack* of broken spatial symmetry — in order to get a zero charge current — and a finite RSOI ensuring a breaking of the spin symmetry thus allowing for a finite spin current. Theory predicts spin ratchets to work both in the ballistic regime (Scheid *et al.*, 2007b, 2010; Ang *et al.*, 2015) and in the dissipative one (Smirnov *et al.*, 2008a,b, 2009). A set-up for the ballistic ratchet has been also proposed by using periodically disposed magnetic stripes instead of electrostatic barriers and RSOI (Scheid *et al.*, 2007a, 2006).

A ballistic spin ratchet is mainly constituted by a quasi-one dimensional channel with a set of symmetric barriers and RSOI [see figure 9(a)]. The lack of broken symmetry implies the absence of a net charge current. In order to understand the presence of a finite spin current we show here the same argument proposed in Refs. [Scheid *et al.*, 2007b, 2010]. A key ingredient is

the sub-band hybridisation that we have introduced in Sec. III.A.

We consider a wire with two open transverse modes ( $N = 1, 2$ ) and a smooth symmetric potential barrier  $U(x)$  in the two bias or *rocking* situations ( $V \gtrless 0$ ), see figure 9(b). Upon adiabatically traversing the barrier from

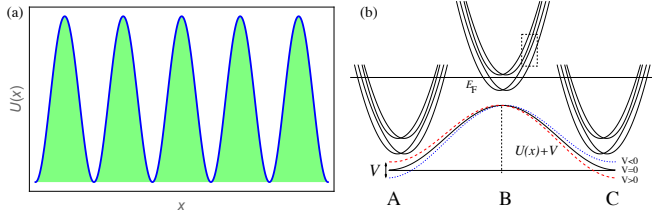


FIG. 9 (Colours online) Panel (a): Sketch of the system of symmetric periodic equally spaced barriers used for the ballistic spin ratchet. Panel (b): Illustration of the spin polarisation mechanism for transmission through a strip with a single adiabatic symmetric potential barrier  $U(x)$  (solid line) in the two rocking situations (dashed and dotted line). At points A, B and C the position-dependent energy dispersion relation  $E_n(k_x)$  is sketched with respect to the Fermi energy  $E_F$  (horizontal line) for two transverse modes and RSOI-induced spin splitting of each mode. Courtesy of Ref. [Scheid *et al.*, 2007b]

region A via B to C [see lower part of figure 9(b)], the electron energy spectrum  $E_n(k_x)$  split by RSOI is shifted rigidly upward and downward by the adiabatic potential barrier  $U(x)$ . For fixed Fermi energy  $E_F$ , the initial shift causes a depopulation of the upper levels ( $N = 2$ ) and a spin-dependent repopulation of the lower one ( $N = 1$ ) while moving from B to C. If  $E_F$  is traversing an anti-crossing between successive modes (see the region indicated by the dashed window in figure 9(b)), there is a certain probability  $P$  for electrons to change their spin state. This causes an asymmetry between spin-up and spin-down states for the repopulated levels (Eto Mikio *et al.*, 2005). The related transition probability can be computed in a Landau-Zener framework. In the case of a transverse parabolic confinement of frequency  $\omega_0$ , this probability reads (Eto Mikio *et al.*, 2005; Scheid *et al.*, 2007b):

$$P(\pm V_0) = 1 - \exp \left\{ \frac{-\pi k_{\text{SO}} \omega_0 \Sigma_z^{-1}}{\partial_x [U(x, z) \pm V_0 g(x, z)]} \right\}. \quad (42)$$

Here  $\Sigma_z$  denotes the difference in the polarisation of the two modes involved, and the function  $g(x, z)$  describes how the potential drops inside the system. In principle  $g(x, z)$  is obtained by solving the Boltzmann equation, however its exact form is irrelevant in order to understand the working principle of the spin ratchet. The spin transmission is proportional to  $P(V)$  and thus different for the two rocking situations  $V \gtrless 0$ . Hence, the ratchet spin current ( $I_S(V)$ ) is finite, also for the case of a symmetric barrier. A quantitative explanation of the spin ratchet effect for a system of multiple barriers [figure 9(a)] is beyond this model. An experimental verification of the spin

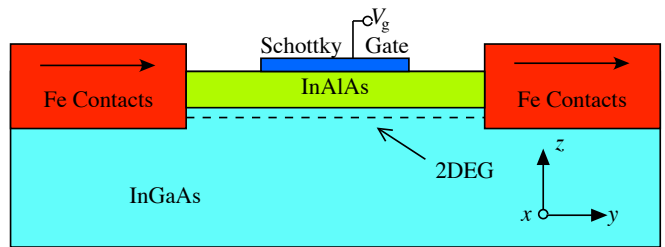


FIG. 10 (Colours online) The spin-field effect transistor proposed by Datta and Das [Datta and Das, 1990].

ratchet effect was proposed in Ref. [Costache and Valenzuela, 2010]. However, in this experiment the breaking of the spin symmetry was achieved by combining a superconductor with a magnetic field.

## 2. Spin Pumping

Adiabatic charge pumping (Brouwer, 1998) consists of the transport of charge obtained — at zero bias voltage — through the periodic modulation of some parameters in the scattering region, *e.g.* some voltages.

In 2003, Governale *et al.* [Governale *et al.*, 2003] proposed an updated of the scheme of charge pumping extended to the spin, following a suggestion by Mucciolo *et al.* [Mucciolo *et al.*, 2002]. The pumping scheme consists of an electrostatic barrier that is changed periodically in time and a gate, as second pumping parameter, to tune the strength of RSOI (Governale *et al.*, 2003). They showed that, neglecting sub-band hybridization, the charge current is zero whereas the spin current is finite. However, in a more realistic case in which many modes are opened, and consequently coupled by RSOI, the charge current is finite but two order of magnitude smaller than the spin one. A prototypal spin pumping system has been realised in a GaAs quantum dot in Ref. [Watson *et al.*, 2003], in which an ingenious method for detecting spin currents is presented. Similar pumping mechanisms have been proposed for graphene with RSOI (Bercieux *et al.*, 2012), and TIs (Citro *et al.*, 2011; Inhofer and Bercieux, 2013; Hofer and Büttiker, 2013; Hofer *et al.*, 2014).

## B. Datta and Das spin Field Effect Transistor

The spin momentum locking due to the RSOI is one of the key ingredients used in 1990 for the proposal of a spin-FET by Datta and Das [Datta and Das, 1990]. It is believed that spin-FET has the advantages of low energy consumption and fast switching speed since it does not involve creating or eliminating the electrical conducting channels during the switching, required by traditional FETs. This spin-FET is the electronic analog

of an electro-optic modulator. The electro-optical effect makes the dielectric constant of a medium different along the two perpendicular directions (*e.g.*,  $y, z$ ).

Let us assume to have photons with a polarisation at  $45^\circ$  with respect to the  $y$  axis (in the  $y$ - $z$  plane) (Born and Wolf, 1999). They can be represented as a linear combination of  $z$ - and  $y$ -polarised photons:

$$\begin{pmatrix} 1 \\ 1 \end{pmatrix}_{45^\circ} = \begin{pmatrix} 1 \\ 0 \end{pmatrix}_z + \begin{pmatrix} 0 \\ 1 \end{pmatrix}_y. \quad (43)$$

Because of the anisotropy of the dielectric constant, as the light passes through the electro-optic material of length  $L$ , the two components acquire different phase shifts  $k_1 L$  and  $k_2 L$ . Thus the light emerging from the electro-optic material can be represented as  $(e^{ik_1 L}, e^{ik_2 L})^T$ . It is collected by an analyser polarised along the  $45^\circ$  direction. Therefore the output power is given by

$$P_0 \propto \left| (1 \ 1) \begin{pmatrix} e^{ik_1 L} \\ e^{ik_2 L} \end{pmatrix} \right|^2 = 4 \cos^2 \frac{(k_1 - k_2)L}{2}. \quad (44)$$

The light output is modulated with a gate voltage that controls the differential phase shift  $\Delta\theta = (k_1 - k_2)L$ .

The analog electronic device based on the RSOI is shown in figure 10. In the original proposal by Datta and Das, the polariser and the analyser were suggested to be implemented via ferromagnetic contacts (as Fe) (Johnson and Silsbee, 1988). In such materials the density of states for electrons with a specific spin orientation — at the Fermi energy — greatly exceeds that for the opposite direction. A contact magnetised in the  $x$  direction preferentially injects and detects electrons spin polarised along positive  $x$  which is represented as a linear combination of the positive  $z$ -polarised  $|\uparrow\rangle$  and negative  $z$ -polarised electrons  $|\downarrow\rangle$ :

$$\begin{pmatrix} 1 \\ 1 \end{pmatrix}_x = \begin{pmatrix} 1 \\ 0 \end{pmatrix}_z + \begin{pmatrix} 0 \\ 1 \end{pmatrix}_z \quad (45)$$

Finally, the analogue of the electro-optic material is realised by employing a 1D quantum wire with RSOI. Within a single band model, RSOI causes the  $|\uparrow\rangle$  and  $|\downarrow\rangle$  electrons with the same energy to have different wave vectors  $k_\uparrow$  and  $k_\downarrow$  [see figure 5(e)]. Let us consider an electron travelling in the  $x$  direction with  $k_z = 0$  and  $k_x \neq 0$ . The eigenenergies for the corresponding 1D case are given equation 9 with  $k_y = 0$ .

Using equations (9)-(14), it is possible to recover a relation for the phase shift between the two spins:

$$\Delta\theta = (k_\uparrow - k_\downarrow)L = \frac{2m\alpha L}{\hbar^2} \quad (46)$$

which is proportional to  $\alpha$ .

The above analysis was originally limited to a single-mode quantum wire. However, as we have learnt in the

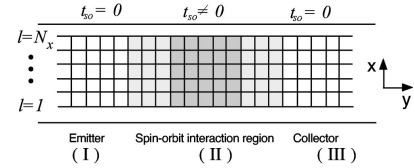


FIG. 11 Schematic of the tight-binding model for the system. In the shaded areas the spin-orbit interaction if finite  $t_{SO} \neq 0$ . Courtesy of Ref. [Mireles and Kirczenow, 2001].

Sec. III.B, in the presence of multiple modes, physics changes.

A fully numerical multi-mode analysis of the spin-FET was proposed by Mireles and Kirczenow (Mireles and Kirczenow, 2001). They investigate the effect of the strength of the RSOI on the spin-transport properties of narrow quantum wires of width  $W$ . This is a quasi-one-dimensional wire, *i.e.*, assumed to be infinitely long in the propagation direction (see figure 11).

The nearest-neighbour tight-binding Hamiltonian the RSOI (5) takes the form

$$\mathcal{H}_{SO}^{\text{tb}}(y) = -t_{SO} \sum_{s,s'} \sum_{\langle \ell, \ell' \rangle_y} \sum_{\langle j, j' \rangle_x} \left( c_{\ell, m, s'}^\dagger (is_y)_{s, s'} c_{\ell', m, s} - c_{\ell, j, s'}^\dagger (is_x)_{s, s'} c_{\ell', j', s} \right) \quad (47)$$

where the summation is intended over next-neighbours  $\langle \dots \rangle_{x,y}$ ,  $t_{SO} = \frac{a}{2} a$  is the isotropic next-neighbour transfer RSOI hopping term (where  $a$  is the lattice constant), and  $c_{\ell, j, s}^\dagger$  the electron creation operator on the site  $(\ell, j)$  with spin state  $s$  ( $s = \uparrow, \downarrow$ ).

The wire is divided in three main regions. In two of these [I and III in figure 11], which are near the ferromagnetic source and drain, the parameter  $t_{SO}$  is set to zero.

In the middle region (II) the RSOI is finite ( $t_{SO} \neq 0$ ).

The spin-dependent transport problem is solved numerically through the use of the spin-dependent

Lippman-Schwinger equation<sup>4</sup>

$$|\Psi\rangle = |\Phi\rangle + G_0(E)\hat{\mathcal{H}}_{\text{SO}}^{\text{tb}}|\Psi\rangle, \quad (48)$$

where  $|\Phi\rangle$  is the unperturbed wave function, *i.e.* an eigenstate of the kinetic lattice Hamiltonian  $\mathcal{H}_0$  without RSOI, and  $G_0(E) = (E + i\epsilon - \mathcal{H}_0)^{-1}$  is the Green's functions for the system in the absence of any kind of scattering.

A simple criterion for distinguishing the cases of *weak* and *strong* RSOI is introduced in Ref. (Mireles and Kirczenow, 2001). The contribution of the mixing of the spin sub-bands should be negligible as long the sub-bands spacing  $\Delta E_W = E_m^0 - E_n^0$  is much larger than the sub-band intermixing energy

$$\frac{\langle \phi_{n,s} | \mathcal{H}_{\text{SOI}} | \phi_{m,s'} \rangle}{E_m^0 - E_n^0} \ll 1 \quad (49)$$

where  $|\phi_{n,s}\rangle$  are the unperturbed electron wavefunctions. However, if the confinement energy and/or the RSOI are of the same order as the energy shift introduced by the sub-band mixing contribution, then the above condition is about one or larger. In this case one can introduce

$$\beta_{\text{SO}} \approx \frac{(\frac{\pi a}{W})^2}{(\frac{\pi a}{W}) + ak_F} = \beta_{\text{SO}}^c, \quad (50)$$

where  $\beta_{\text{SO}} = \frac{t_{\text{SO}}}{|t|}$ , and  $k_F$  is the Fermi wave number. The critical value  $\beta_{\text{SO}}^c$  defines a *weak* RSOI regime if  $\beta_{\text{SO}} < \beta_{\text{SO}}^c$  and a *strong* coupling regime if  $\beta_{\text{SO}} > \beta_{\text{SO}}^c$  (Mireles and Kirczenow, 2001).

In figure 12 the behaviour of the spin-resolved conductance as function of the RSOI parameter  $t_{\text{SO}}$  is shown. In figure 12(a) the incident Fermi energy is fixed to 0.5 ( $k \approx 0.7a^{-1}$ ) and  $W = 6a = 60$  nm, which gives a critical value  $\beta_{\text{SO}}^c = 0.22$ . This value of  $\beta_{\text{SO}}$  separates the sinusoidal behaviour of  $G^{\uparrow\downarrow}$  for  $\beta_{\text{SO}} \leq 0.22$  from its behaviour for  $\beta_{\text{SO}} > 0.22$  where the confinement energy is of the order of the sub-band mixing energy. The effect is clearer for a wider wire ( $W = 120$  nm) [see figure 12(c)] for which the critical value of  $\beta_{\text{SO}}$  is 0.07. To show that the non-sinusoidal behaviour is due mainly to the sub-band mixing, in the figure 12(b) and 12(d) we report the spin-conductance as function of  $t_{\text{SO}}$  with the same parameter of figure 12(a) and 12(c) respectively but in the unphysical situation of  $\alpha_y \neq 0$  and  $\alpha_x = 0$ , respectively. It is evident that the sinusoidal behaviour is recovered. The modification of spin-FET due to the inclusions of many transversal modes has been also investigated in Ref. [Jeong and Lee, 2006].

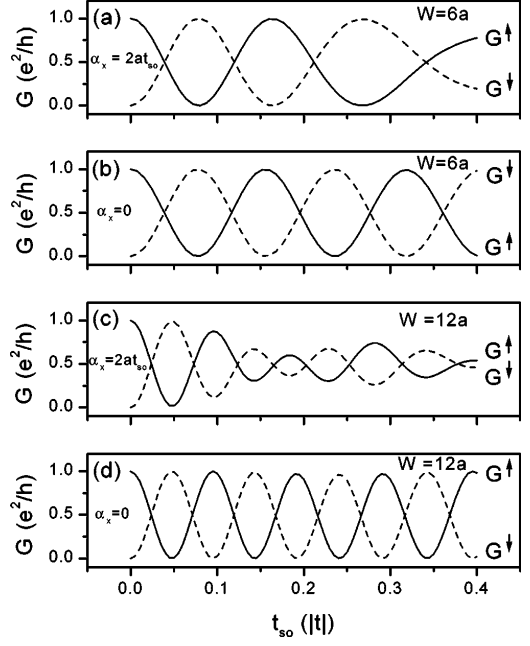


FIG. 12 Spin-orbit coupling strength dependence of the ballistic spin conductance; solid line is  $G^{\uparrow}$ , dashed line  $G^{\downarrow}$ : (a) Narrow wire of  $W = 6a$  and uniform RSOI ( $\alpha_x = \alpha_y = 2at_{\text{SO}}$ ). (b) Same as in (a) but with  $\alpha_x = 0$  and  $\alpha_y = 2at_{\text{SO}}$ ; perfect oscillations are seen for all  $t_{\text{SO}}$ . (c) Same as in (a) with  $W = 12a$ . (d) Modulation for  $W = 12a$ , with  $\alpha_x = 0$  and  $\alpha_y = 2at_{\text{SO}}$ . The sub-band mixing clearly changes the otherwise perfectly sinusoidal spin-conductance modulation. Courtesy of Ref. [Mireles and Kirczenow, 2001].

So far, several obstacles have been found on the way of the realisation of the spin-FET (Datta and Das, 1990). The main one is related to the injection of spin polarised currents. For example, it has been shown that in diffusive transport regime, for typical ferromagnets only a current with a small polarisation can be injected into a semiconductor 2DEG with long spin-flip length even if the conductivity of semiconductor and ferromagnet are equal (Schmidt *et al.*, 2000). This situation is dramatically exacerbated when ferromagnetic metals are used; in this case the spin polarisation in the semiconductor is negligible.

A possible solution to this problem was proposed by employing dilute magnetic semiconductor (Ohno, 1998) as source and drain contacts. In these systems a few percent of the cations in the III-V or II-VI semiconductors compounds are randomly substituted by magnetic ions, usually Mn, which have local magnetic moments. The effective coupling between these local moments is mediated by free carriers in the host semiconductor compound (holes for *p*-doped materials and electrons for *n*-doped one) and can lead to ferromagnetic long-range order. Curie temperatures  $T_c$  close to 100 K have been found in bulk (Ga,Mn)As systems (Ohno, 1998).

<sup>4</sup> Nowadays a lot of numerical computation, also for systems with RSOI, is performed via recursive Green's function method; among the free codes one of the most used is KWANT (Groth *et al.*, 2014).



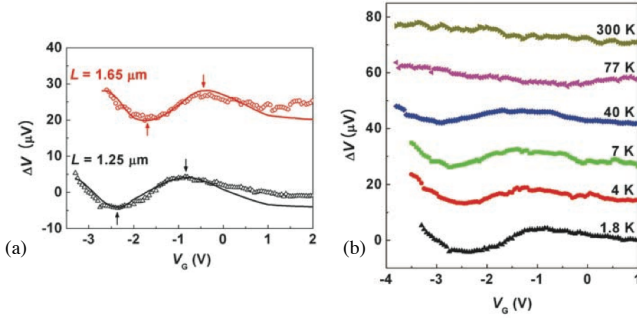


FIG. 13 (Colours online) (a) Gate voltage modulation of spin FETs having different channel lengths, with  $T = 1.8$  K and  $I = 1$  mA. The symbols indicate experimental data. The solid lines are the fits obtained from  $V = A \cos(2m^* \alpha L \hbar^{-1} + \varphi)$ . Data are offset for clarity. Baseline voltages are 1.032 mV and 0.715 mV for  $L = 1.25 \mu\text{m}$  and  $1.65 \mu\text{m}$ , respectively; (b) Temperature dependence of oscillatory conductance with  $L = 1.25 \mu\text{m}$  and  $I = 1$  mA. As temperature increases, the mean free path decreases and transport characteristics change from ballistic to diffusive. Courtesy of Ref. [Koo *et al.*, 2009].

Using the properties of the dilute magnetic semiconductor, all-semiconductor spin-FET have been proposed in which the conducting channel is provided by a two-dimensional hole gas (Pala *et al.*, 2004).

The first experimental realisation of the spin-FET has been done in 2009 by the Johnson's group (Koo *et al.*, 2009). Instead of using the configuration initially suggested by Datta and Das (Datta and Das, 1990), in this experiment a nonlocal measurement scheme has been used (Johnson and Silsbee, 1988; Jedema *et al.*, 2001, 2002; Lou *et al.*, 2007). This allows to perform a better measurement of the charge and spin signals. The quantum wire was realised in a InAs heterostructure with strong RSOI with the ferromagnetic contacts realised with  $\text{Ni}_{81}\text{Fe}_{19}$  permalloy on the top. Figure 13(a) shows the oscillation of the output voltage of the spin-FET as a function of the gate voltage that is changing the strength of the RSOI for two different length of the distance between the two ferromagnetic contacts. As reported in equation (46) the phase shift between the two spin channels is proportional to the distance between the two ferromagnets. Therefore, for the longer case (red-solid data) we observe a shift of the length on the half period of oscillations. Figure 13(b) shows the oscillation for a fixed distance between the ferromagnetic contacts but at different temperatures. We observe a signature of coherent oscillations up to 40 K. At higher temperatures inelastic scattering become more pronounced and coherent effects are washed out (Koo *et al.*, 2009). Another experimental realization was also lately reported in Ref. [Chuang *et al.*, 2014].

## V. INTERFERENCE EFFECTS AND BERRY PHASE

### A. Rashba interaction as SU(2) gauge field: application to quantum networks

In this section we explore RSOI as a SU(2) non-Abelian gauge field. We start by recasting the RSOI in the Hamiltonian (8) as a SU(2) vector potential:

$$\begin{aligned} \mathcal{H}_0 &= \frac{1}{2m} \left[ \mathbf{p} + \frac{m^* \alpha}{\hbar} (\mathbf{z} \times \mathbf{s}) \right]^2 - \left( \frac{m^* \alpha}{\hbar} \right)^2 \\ &= \frac{1}{2m^*} (\mathbf{p} + \mathcal{A}_s)^2 - \left( \frac{m^* \alpha}{\hbar} \right)^2 \end{aligned} \quad (51)$$

comparing the two lines of (51) we recognise  $\mathcal{A}_s = \frac{m^* \alpha}{\hbar} (\mathbf{z} \times \mathbf{s})$ . In this form the Rashba Hamiltonian resembles that of a particle in a magnetic field, thus allowing for a straightforward connection with the physics of the AB effect (Aharonov and Bohm, 1959). In the case of a SU(2) vector potential, it is known as AC effect (Aharonov and Casher, 1984). In analogy to AB effect, we can introduce a phase field that reads:

$$\psi_{AC} \equiv \frac{2\pi}{\phi_0} \oint \mathcal{A}_s \cdot d\mathbf{r} = 2\pi \frac{\phi_{\text{SOI}}}{\phi_0}, \quad (52)$$

where  $\phi_{\text{SOI}}$  is the flux associated to the RSOI effective field and  $\phi_0 = \frac{hc}{e}$  the flux quantum. When travelling through a closed path an electron gains a non-Abelian phase due to the presence of the RSOI. This extra phase can give rise to interference phenomena. By contrast to the phase gained in a perpendicular magnetic field — that depends only on the area enclosed by the particle path — here the actual path covered by the electron plays an important role. In the standard AB set-up the magnetic field can be tuned in order to move from a full destructive to a full constructive interference. In the case of AC the role of the magnetic field is played by the RSOI that can be modulated modifying  $\alpha$ , *i.e.* by gating the heterostructure. The AC effect has been observed in semiconductor heterostructure (Bergsten *et al.*, 2006), in HgTe rings (König *et al.*, 2006) and TI interferometric structures (Qu *et al.*, 2011) (c.f. figure 4). The interplay between the AB and the AC effect, and the mutual effect of an Abelian and non-Abelian gauge field has been investigated in Ref. [Nagasawa *et al.*, 2012, 2013].

In order to understand the fundamental difference between the AB and the AC effect, we propose here a very simple *gedankenexperiment* for a square interferometer. Let  $\mathcal{R}_{pq}$  be the phase gained by the wave function  $\Psi(\mathbf{r})$  — travelling from a point  $p$  to a point  $q$ . It reads

$$\mathcal{R}_{pq}^{\text{SU}(2)} = \exp \left\{ -i \int_p^q \mathbf{s} \cdot (\mathbf{z} \times d\mathbf{l}) k_{\text{SO}} \right\} \quad (53a)$$

$$\mathcal{R}_{pq}^{\text{U}(1)} = \exp \left\{ -i \frac{2\pi}{\phi_0} \int_p^q \mathcal{A} \cdot d\mathbf{l} \right\} \quad (53b)$$

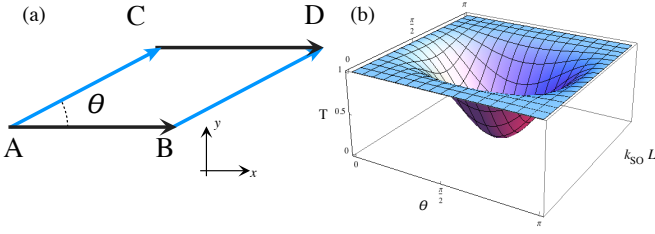


FIG. 14 (Colours online) (a): Scheme of the interferometer for addressing the difference between an Abelian and a non-Abelian phase. (b): Normalised transmission probability  $T \propto \text{Tr}[\Gamma\Gamma^\dagger]$  for the interferometer in the left panel in the case of the action of a  $SU(2)$  gauge field (53a).

depending on whether we are considering a  $U(1)$  or  $SU(2)$  gauge field, where  $\mathcal{A}$  is the vector potential associated to a magnetic field  $\mathbf{B}$ . We now consider the closed path in figure 14(a) and explicitly consider the phases gained by the electron travelling along each of its arm. The four arms of the interferometer have the same length  $\ell_{pq} = L$  ( $pq \in \{AB, AC, BD, CD\}$ ), the angle at its base is  $\theta$ , and its area is  $S$ . In order to obtain destructive interference the sum of the phases gained along consecutive arms has to be zero:

$$\Gamma = \mathcal{R}_{BD}^\beta \cdot \mathcal{R}_{AB}^\beta + \mathcal{R}_{CD}^\beta \cdot \mathcal{R}_{AC}^\beta = 0 \quad (54)$$

with  $\beta \in \{SU(2), U(1)\}$ .

By using the Gauss' theorem and defining the magnetic field flux as  $\phi = |\mathbf{B}|S = (\nabla \times \mathcal{A})S$ , we obtain that in the case of a  $U(1)$  phase the destructive interference is verified when the condition  $\frac{\phi}{\phi_0} = \frac{1}{2}$  is met for every the value of the angle  $\theta$ . Here, the condition for the destructive interference is straightforward because the phases (53b) are c-numbers.

In the case of the non-Abelian phase field the phases (53a) are linear combination of Pauli matrices. Therefore a solution for the condition (54) is obtained by looking at the transmission probability — proportional (to lowest order) to  $\text{Tr}[\Gamma\Gamma^\dagger]$ . This quantity is shown in the figure 14(b), we can see that the condition of zero transmission probability is obtained if  $\theta = \frac{\pi}{2}$  and  $k_{SO}L = \frac{\pi}{2}$ . To summarise, while in the case of a  $U(1)$  gauge field we can obtain destructive interference for each geometrical realisation of the interferometer [see figure 14(a)], in the  $SU(2)$  case a specific geometric arrangement of the loop is required.

These effects can have interesting implications in the transport properties of nontrivial quantum networks. In the following, we shall focus on a very special type of lattice structure: the  $\mathcal{T}_3$  lattice [c.f. figure 15(b)], whose unit cell contains three inequivalent sites (Sutherland, 1986). Its spectrum is composed of two dispersive bands that are equivalent to the one of the honeycomb lattice and a flat band at zero energy that is due to the presence of lattice sites with uneven coordination number (Bercieux et al.,

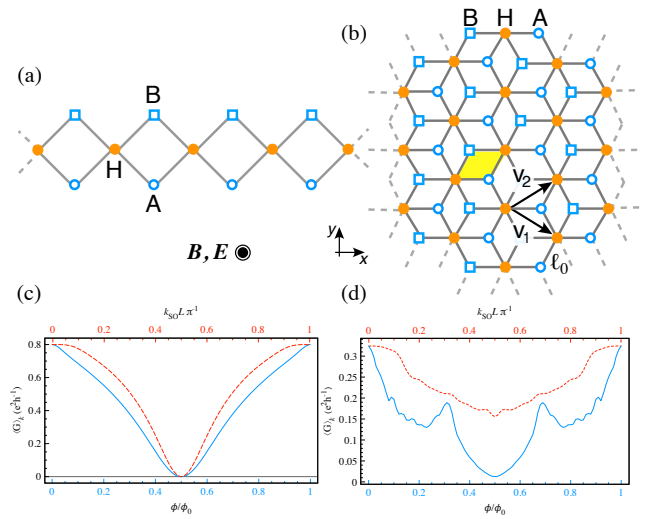


FIG. 15 (Colours online) (a) Sketch of diamond chain, this is the one-dimensional version of the  $\mathcal{T}_3$  lattice. (b): Sketch of the  $\mathcal{T}_3$  lattice, it is a Bravais lattice with a unit cell containing three inequivalent sites: A, B and H, the latter has coordination number higher than the other two (Sutherland, 1986). (c): averaged conductance for the diamond chain as a function of the magnetic flux ratio (lower  $x$ -axis, blue solid-line) and as a function of the dimensionless RSOI (upper  $x$ -axis, red-dashed line) (Courtesy of Ref. [Bercieux et al., 2005b]). (d): averaged conductance normalised to the number of leads for the  $\mathcal{T}_3$  lattice as a function of the magnetic flux ratio (lower  $x$ -axis, blue solid-line) and as a function of the dimensionless RSOI (upper  $x$ -axis, red-dashed line) (Courtesy of Ref. [Bercieux et al., 2005b]). In (c) and (d)  $\mathbf{B}$  and  $\mathbf{E}$  are the applied magnetic field and the electric field, respectively.

2009, 2011). In the presence of a perpendicular magnetic field — when the ratio  $\frac{\phi}{\phi_0} = \frac{1}{2}$  — the spectrum reduces to three flat bands (Vidal et al., 1998). This localisation effect is a consequence of the AB effect taking place within each of the plaquette contained in the  $\mathcal{T}_3$  lattice [in yellow in figure 15(b)]. The same effect is observed in its quasi-1D analog, the diamond chain [c.f. figure 15(a)] for the  $\frac{\phi}{\phi_0} = \frac{1}{2}$  (Vidal et al., 2000a).

Can the same physics be driven also by the RSOI? The question has been addressed by one of the authors using the method of quantum networks (Naud et al., 2001). Full localisation due RSOI is possible only in the diamond chain (Bercieux et al., 2004) [figure 15(c)] and is forbidden, for the arguments illustrated above, in the case of the  $\mathcal{T}_3$  lattice (Bercieux et al., 2005b) — figure 15(d). In figure 15(c) and 15(d) we show the conductance of the diamond lattice and the  $\mathcal{T}_3$  lattice, respectively, as a function of the magnetic flux and RSOI. We observe that in the case of the diamond chain 15(c), both effects induce a complete localisation equivalent to zero conductance. However, in the case of the  $\mathcal{T}_3$  lattice 15(d), only the magnetic field can induce (almost) complete localisation. The residual conductance comes from the con-

ducting state along the lattice boundary due to the Hall effect. However it can be reduced to zero by injecting electrons through the centre of the lattice (Vidal *et al.*, 2000b). Signatures of this AB caging effect have been observed also experimentally (Naud *et al.*, 2001; Abilio *et al.*, 1999). The diamond chain lattice model with RSOI and AB field has been also proposed as a system for producing a spin filter (Hatano *et al.*, 2007; Aharony *et al.*, 2008, 2011; Matityahu *et al.*, 2013a,b).

## B. Interference in quantum rings

Mesoscopic quantum rings allow to have direct access to the phase of the electron wavefunction, when their size is smaller than the coherence length. Interference effects have been observed in metal quantum rings many years ago (Washburn and Webb, 1992). Since electrons are spinful particles, the spin part of the wavefunction is influenced by the magnetic field via the Zeeman term in the Hamiltonian. A more subtle effect arises when there is a magnetic field non-orthogonal to the plane of the orbiting particle (*e.g.* the effective magnetic field due to the RSOI) because, as a consequence of the orbital motion, its spin dynamics is instantaneously governed by a time-dependent Hamiltonian (Anandan, 2002; Saarikoski *et al.*, 2014). This time dependence ends up in an extra phase acquired by the particle wavefunction which is named after Berry [Berry, 1984 and Aharonov and Anandan, 1987], who put in foreground its topological properties when the orbits are closed.

During the last years, the effects of RSOI on the AB oscillations have been observed in semiconductor based quantum rings by several groups (Yau *et al.*, 2002; Nitta *et al.*, 2003; Meijer *et al.*, 2004; Morpurgo *et al.*, 1998). As said before, in the presence of both orthogonal magnetic field and RSOI, the total effective momentum dependent magnetic field is tilted with respect to the vertical direction. The resulting Berry phase influences the interference pattern.

In Refs. [Capozza *et al.*, 2005 and Lucignano *et al.*, 2007] one of the authors studied the conductance and the spin transport in a quantum ring in the presence of RSOI and magnetic field, accounting also for dephasing at the contacts. The AB resonance in the Fourier transform of the magneto-conductance displays satellite peaks due to the RSOI (see figure 16) that have been experimentally observed (Habib *et al.*, 2007). During the last years, several theoretical techniques have been employed to study quantum rings. In Refs. [Loss *et al.*, 1990 and Engel and Loss, 2000], an imaginary time path integral approach is developed to study the conductance of a strictly 1D quantum ring, and its conductance fluctuations in the diffusive limit. In Ref. [Tserkovnyak and Brataas, 2007] a real time path integral approach is applied in the limit of negligible Zeeman splitting. Several research articles discussed the

conductance properties and the spin dependent transport of quantum rings in the 1D ballistic limit, by means of a spin dependent scattering matrix approach (Frustaglia and Richter, 2004; Bercieux *et al.*, 2005a; Molnár *et al.*, 2004; Citro *et al.*, 2006; Ramaglia *et al.*, 2006). In the absence of the magnetic flux, the conductance shows quasi-periodic oscillations in the RSOI strength, which can be modified by switching the magnetic field on. Numerical calculations (Frustaglia and Richter, 2004; Souma and Nikolić, 2004; Wu and Cao, 2006) have shown that in the 2D case there are only quantitative modifications of the 1D results that do not qualitatively affect the physics. Thus, in the following, we will just focus on the 1D limit.

The model Hamiltonian, describing a half integer spin particle, in an orthogonal magnetic field and with RSOI (Meijer *et al.*, 2002), reads

$$\mathcal{H}[\mathbf{p}, \mathbf{r}, \mathbf{S}] = \frac{1}{2m} \left( \mathbf{p} + \frac{e}{c} \mathbf{A}_0 \right)^2 - \omega_c S_z + \mathcal{H}_{\text{RSOI}} \quad (55)$$

$$\mathcal{H}_{\text{RSOI}} = \frac{2\alpha}{\hbar^2} \left[ \mathbf{z} \times \left( \mathbf{p} + \frac{e}{c} \mathbf{A}_0 \right) \right] \cdot \mathbf{S},$$

where  $\mathbf{A}_0(\mathbf{r}) = \frac{B}{2}(-y, x, 0)$  is the vector potential generating the uniform field  $B$ , normal to the ring surface, and  $\omega_c = \frac{q^* e B}{2mc}$  is the cyclotron frequency. We fix the vector potential in the symmetric gauge. We present, here, fully general results as they depend on the ratio  $\frac{\alpha}{\hbar\omega_c R}$  which can be tuned by acting on  $\alpha$ . We assume a single channel ring as a 1D circle of radius  $R$ , connected to two leads. Accordingly, the position of the particle on the ring is parametrised by the angle  $\varphi$ . The vector potential has just the azimuthal component  $A_\varphi = \frac{\phi}{2\pi R}$ , where  $\phi$  is the magnetic flux threading the ring.

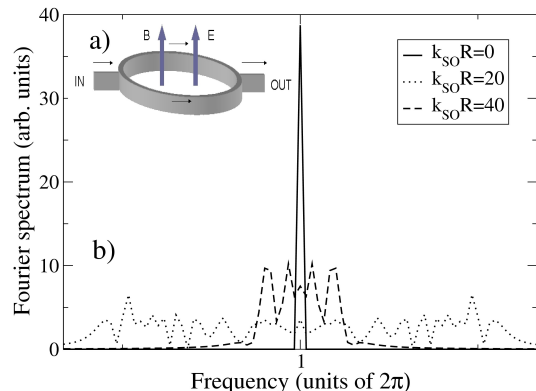


FIG. 16 (Colours online) a): Sketch of a quantum ring. The magnetic field  $B$  drives the AB effect, while the electric field can tune the RSOI. b) Fourier transform of the magneto-conductance. By increasing ROI two satellite peaks rise close to the AB peak. They can be attributed to the Berry phase acquired by the electron during its motion. Courtesy of Ref. [Capozza *et al.*, 2005].

In order to study the conduction properties of the ring, we need the propagation amplitude for an electron at energy  $E_0$  entering the ring with spin polarisation  $s_0$  and leaving it with spin polarisation  $s_f$ . This is given by

$$A(s_f; s_0 | E_0) = \int_0^\infty \frac{dt_f}{\tau_0} e^{i \frac{E_0 t_f}{\hbar}} \langle \mathbf{r}_f, s_f, t_f | \mathbf{r}_0, s_0, t_0 \rangle, \quad (56)$$

where  $\langle \mathbf{r}_f, s_f, t_f | \mathbf{r}_0, s_0, t_0 \rangle$  is the amplitude for a particle entering the ring at the point  $\mathbf{r}_0$  and at the time  $t_0$  with spin polarisation  $s_0$  to exit at the point  $\mathbf{r}_f$  at the time  $t_f$  with spin polarisation  $s_f$ . Here,  $\tau_0 = \frac{mR^2}{2\hbar}$  is the time scale for the quantum motion. In order to compute  $\langle \mathbf{r}_f, s_f, t_f | \mathbf{r}_0, s_0, t_0 \rangle$ , we adopt a path integral representation for the orbital part of the amplitude. Since we parametrise the orbital motion of the particle in terms of the angle  $\varphi$ , we provide the appropriate Lagrangian,  $\mathcal{L}_{\text{orb}}$ , as a function of  $\varphi, \dot{\varphi}$ . It reads

$$\begin{aligned} \mathcal{L}_{\text{orb}}[\varphi(t), \dot{\varphi}(t), \boldsymbol{\sigma}] = & \frac{m}{2} R^2 \dot{\varphi}^2(t) - \frac{\phi}{\phi_0} \hbar \dot{\varphi}(t) \\ & + \frac{\alpha^2 m}{2\hbar^2} + \frac{\hbar^2}{8mR^2}. \end{aligned} \quad (57)$$

The last two contributions in equation (57) are constants that come from the RSOI term, and the Arthurs term, which is required when a path integration is performed in cylindrical coordinates. Since both contributions are constant, they can be lumped into the incoming energy  $E_0 \sim E_F$  and therefore they will be omitted henceforth.

By taking into account the spin degree of freedom, as well, we represent the propagation amplitude as

$$\begin{aligned} \langle \mathbf{r}_f, s_f, t_f | \mathbf{r}_0, s_0, t_0 \rangle &= \langle \mathbf{r}_f, s_f | e^{-i \int_{t_0}^{t_f} dt \mathcal{H}} | \mathbf{r}_0, s_0, t_0 \rangle \\ &= \int_{\varphi(t_0)=\varphi_0}^{\varphi(t_f)=\varphi_f} \mathcal{D}\varphi e^{-i \int_0^{t_f} dt [\tau_0 \dot{\varphi}^2 - q\dot{\varphi}]} \langle s_f | \mathbf{U}_{\text{spin}}(t_f, t_0) | s_0 \rangle, \end{aligned} \quad (58)$$

where  $q = \frac{\phi}{\phi_0}$ .

$$\mathbf{U}_{\text{spin}}(t_f, t_0) = \mathbf{T} \exp \left[ -\frac{i}{\hbar} \int_{t_0}^{t_f} d\tau \mathbf{H}_{\text{spin}}(\tau) \right]. \quad (59)$$

is the full spin propagator,  $\mathbf{T}$  is the time ordering operator, and  $\mathbf{H}_{\text{spin}}(t)$  is the spin Hamiltonian that reads

$$\mathbf{H}_{\text{spin}}(t) = \frac{\hbar}{2} \begin{pmatrix} \omega_c & 2\gamma\dot{\varphi}e^{-i\varphi(t)} \\ 2\gamma\dot{\varphi}e^{i\varphi(t)} & -\omega_c \end{pmatrix}, \quad (60)$$

with  $\gamma = \frac{2\alpha\tau_0}{\hbar R}$ .

For a ring device, at each lead one has to take into account three possible scattering processes, consistently with the conservation of the total current. This is described in terms of a unitary  $\mathcal{S}$ -matrix that, when the two arms are symmetric, is given by

$$\mathcal{S} = \begin{pmatrix} -\frac{1}{2}(1+r) & \frac{1}{2}(1-r) & \sqrt{\frac{1}{2}(1-r^2)} \\ \frac{1}{2}(1-r) & -\frac{1}{2}(1+r) & \sqrt{\frac{1}{2}(1-r^2)} \\ \sqrt{\frac{1}{2}(1-r^2)} & \sqrt{\frac{1}{2}(1-r^2)} & r \end{pmatrix}. \quad (61)$$

The numerical labelling of the  $\mathcal{S}$ -matrix elements referring to the three terminals of each contact fork, are explained in figure 17(1a). Assuming, for simplicity, that the scattering matrix is the same for both leads, equation (61) will hold both at the left-hand lead, and at the right-hand lead of the ring.

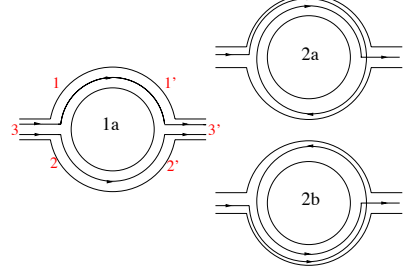


FIG. 17 (Colours online) First and second order paths included in the calculation of the transmission amplitude across the ring, from left to right, including forward scattering only. Numbers 1, 2, 3(1', 2', 3') in figure 17a refer to the labelling of the terminals in Eq.(61). Courtesy of Ref. [Lucignano *et al.*, 2007].

In particular,  $\mathcal{S}_{3,3} = r$  is the reflection amplitude for a wave coming from the left lead,  $\mathcal{S}_{1(2),1(2)} = -\frac{1}{2}(1+r)$  the reflection amplitude for a wave incoming from the upper/lower arm,  $\mathcal{S}_{1(2),2(1)} = \frac{1}{2}(1-r)$  is the transmission amplitude from the upper (lower) to the lower (upper) arm and  $\mathcal{S}_{1(2),3} = \mathcal{S}_{3,1(2)} = \sqrt{\frac{1}{2}(1-r^2)}$  is the transmission amplitude from the upper/lower arm to outside of the ring. As the ring is assumed to be symmetric, the same scattering matrix applies to the right lead where we indicate with primed numbers the three terminals as shown in figure 17. In figure 17 we show the simplest paths of the electrons in the ring including only forward scattering at the contacts. More involuted paths arise if we account also for backscattering processes in which the electron can get backscattered within the same ring's arm from which it is coming. For instance, the paths (2f) and (2h), as well as (2g) and (2i) in figure 18, include looping in opposite directions around the ring. Interference between clockwise and counterclockwise windings leads to WL corrections. We denote these corresponding paths — including also (2c) and (2d) — as “reversed paths”. In our approach, all order paths are numerically generated up to the convergency and the  $\mathcal{S}$ -matrix (61) is implemented in the numerical algorithm. Within Landauer's approach, the conductance  $\mathcal{G}$  is given by

$$\mathcal{G} = \frac{e^2}{\hbar} \sum_{s,s'} |\mathcal{A}(s; s' | E_F)|^2. \quad (62)$$

Here we will consider the dependence on the external magnetic field  $\frac{\phi}{\phi_0}$  and on the RSOI strength  $k_{\text{SO}}R$  both in the absence and in the presence of dephasing at the contacts. To make the model more realistic, we allow for

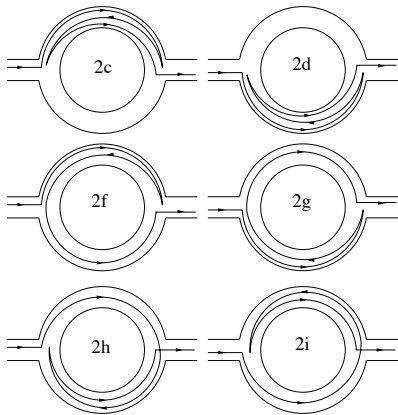


FIG. 18 Second order paths of the transmission amplitude from left to right including backscattering at the leads. Paths (2f) and (2h), as well as (2g) and (2i) contribute to the weak localization corrections. Courtesy of Ref. [Lucignano *et al.*, 2007].

higher order looping of the electron within the ring. In Ref. [Capozza *et al.*, 2005], only the paths of the kind of figures 17(2a) and 17(2b) were included. Following Ref. [Lucignano *et al.*, 2007] we consider here also the paths of the kind of figure 18 in which the electron can be backscattered into the ring. We use here  $r = 0$  in the scattering matrix between the arms and the leads, which means that no back-reflection in the incoming lead is present.

The dephasing due to diffusiveness in the contacts is accounted by adding a random phase  $z \in (-\zeta, \zeta)$  for each scattering at the leads. In figure 19, we report the conductance  $\mathcal{G}$  as a function of  $\frac{\phi}{\phi_0}$ , with  $k_{\text{SO}}R = 0$  (left column) or as a function of  $k_{\text{SO}}R$  with  $\frac{\phi}{\phi_0} = 0$  (right column). These are averaged over  $N = 1000$  realisations of dephasing, and plotted for increasing window of phase randomness ( $\zeta = \pi/3, \pi, 2\pi$  from top to bottom). The black curves refer to ideal contacts (including only the paths of figure 17) while the red curves refer to realistic contacts (including also the paths of figure 18).

The ring is rather insensitive to small dephasing at the contacts however, by increasing the amount of dephasing (middle and bottom left panels in figure 19) we find that the sensitiveness is larger in the case of realistic contacts. This is due to the fact that for realistic coupling, the electrons in the ring can experience higher order paths, since it scatters with the leads many times.

In the right panel of figure 19, we plot the *dc*-conductance as a function of  $k_{\text{SO}}R$  at  $\phi = 0$  for both ideal contacts and realistic contacts (black and red lines in each box), with an increasing phase randomisation (boxes

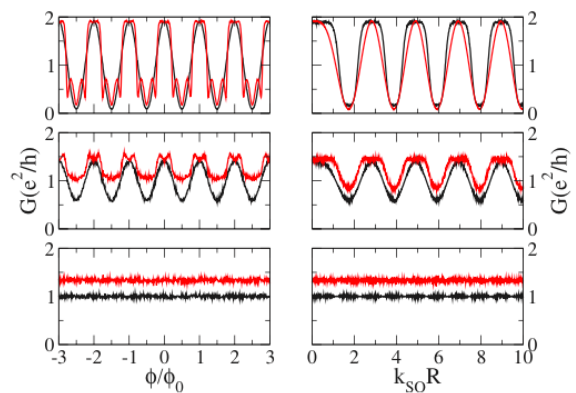


FIG. 19 (Colours online) Conductance as a function of  $\phi/\phi_0$  (left panels) and  $k_{\text{SO}}R$  (right panels) for ideal (black curves) and realistic (red curves) contacts. An increasing amount of dephasing at the contacts is also included: from top to bottom:  $\zeta = \pi/3, \pi, 2\pi$ . Courtesy of Ref. [Lucignano *et al.*, 2007].

from top to bottom with  $\zeta = \pi/3, \pi, 2\pi$ ), averaged over  $N = 1000$  disorder realisations. In the case of ideal contacts and little dephasing (top right panel black curve), we observe again quasi periodic oscillation of the conductance reproducing, the localisation conditions at the expected values of  $k_{\text{SO}}R$  (Frustaglia and Richter, 2004; Bercieux *et al.*, 2005a; Capozza *et al.*, 2005; Lucignano *et al.*, 2007; Molnár *et al.*, 2004). When including higher order processes, interference effects give rise to a slightly different pattern. In the case of realistic contacts, we note that the conductance of the ring is seriously affected by dephasing. Indeed, large dephasing gives rise to random oscillations that are not averaged out — thus washing out the conductance oscillations. The effect takes place for  $\zeta \sim \pi$  when time reversed paths are included, in contrast to  $\zeta \sim 2\pi$  when the time reversed paths are absent. As regular magneto-conductance oscillations are experimentally observed (Yau *et al.*, 2002; Nitta *et al.*, 2003; Meijer *et al.*, 2004; Morpurgo *et al.*, 1998) with little percentage of contrast between maxima and minima, we conclude that, in real samples, dephasing is ubiquitous.

## VI. RELATED PROBLEMS

### A. Spin-Hall effect

The SHE is a phenomenon, associated to SOI — it can be used to electrically generate or detect spin currents in non-magnetic systems. This effect has been observed both in metallic and semiconductor systems (for a review see Ref. [Jungwirth *et al.*, 2012 and Vignale, 2010]). In this review, we will mainly focus on semiconductors, in which RSOI plays a major role. The SHE predicts that an unpolarised electric current can generate a transverse spin current, whose spin is perpendicular to the plane

of the two currents. It was predicted in two pioneering articles by Dyakonov and Perel in 1971 [D'yakonov and Perel, 1971a,b]. It is a consequence of the Mott scattering of electrons on unpolarised impurities, which results in spatial separation of electrons with opposite spins. It is closely related to the anomalous Hall effect — Hall effect in ferromagnetic materials — originally observed by Hall himself [Hall, 1881] and later explained by Karplus and Luttinger [Karplus and Luttinger, 1954] and Nozieres and Lewiner [Nozieres and Lewiner, 1973]. It does not require magnetic field nor magnetism, in other words it does not require broken time reversal symmetry (for a review see [Nagaosa *et al.*, 2010]). The SHE was hardly investigated till Hirsch [Hirsch, 1999] and Zhang [Zhang, 2000] proposed such phenomenon to the attention of the spintronics community. Three main mechanisms have been proposed to contribute to the SHE: namely the spin dependent band structure of the material (the so called intrinsic mechanism) and the impurity scattering mechanisms, *i.e.* the “skew scattering” and the “side jump”, conventionally addressed as extrinsic mechanisms. The extrinsic mechanisms are the same responsible of the anomalous Hall effect (Nagaosa *et al.*, 2010): the skew scattering originates from the different scattering angle due to a spin-orbit impurity, depending on the electron spin and angular momentum; while the side jump reflects the shift in the trajectory of the backscattered electrons originating from the anomalous velocity operator in SOI systems (Crépieux and Bruno, 2001). The intrinsic mechanism was proposed by Murakami *et al.* [Murakami *et al.*, 2003] and by Sinova *et al.* [Sinova *et al.*, 2004], it resorts on the DSOI and RSOI in 2D III-V GaAs quantum wells. This mechanism does not explicitly depend on impurities, however these cannot be neglected as they are absolutely essential to the establishment of the steady state current. In principle, the spin Hall current can be calculated using the Kubo formula. However, inclusion of vertex corrections due to impurities, in the linear SOI model leads to a vanishing spin Hall conductivity (Raimondi and Schwab, 2005; Dimitrova, 2005) in striking contrast to earlier calculations not including vertex corrections (Sinova *et al.*, 2004). More complicated models (explicitly carried out for transition metals) (Tanaka *et al.*, 2008) overtake this problem, giving rise to a finite spin Hall conductivity in reasonable agreement with the experiments. Some proposals require the presence of a magnetic field to give rise to a finite spin Hall current even in the presence of linear SOI and disorder (Lucignano *et al.*, 2008; Engel *et al.*, 2007; Lin *et al.*, 2006; Milletari *et al.*, 2008; Raimondi *et al.*, 2006; Gorini *et al.*, 2008). Among the first successful experiments we here highlight the work by Wunderlich *et al.* [Wunderlich *et al.*, 2005c] that used coplanar *p-n* diodes to detect circularly polarised electroluminescence at opposite edges of the spin Hall bar, and the work by Kato *et al.* [Kato *et al.*, 2004] that employed a magneto-optical Kerr microscope to scan the spin po-

larisation across the Hall bar. The former ascribed their finding to the intrinsic mechanism whereas the latter to the extrinsic one. Since that time several interesting experiments performing optical measurements for the spin detection in the intrinsic and extrinsic SHE have been performed (Matsuzaka *et al.*, 2009; Nomura *et al.*, 2005; Sih *et al.*, 2005b, 2006b; Stern *et al.*, 2006, 2007, 2008; Zhao *et al.*, 2006; Okamoto *et al.*, 2014). A detailed description of this physics is out of the purpose of this reviews and we refer the readers to more specific review articles on the SHE for more information (Jungwirth *et al.*, 2012; Vignale, 2010).

## B. Diffusive limit: weak anti-localization

In the previous sections we have mainly described the physics of ballistic mesoscopic systems, *i.e.* solid state devices in which the lateral size of the sample  $L$  is smaller than the coherence length  $L_\phi$  and of the elastic mean free path  $\ell_e$ . In this regime, impurity scattering can be safely disregarded, and the ballistic motion of electrons well describes the physical scenario. However, in the mesoscopic limit, it may happen that the lateral size of the sample  $L$ , while being still smaller (or of the order of) than the coherent length of the material  $L < L_\phi$ , it is larger than the elastic mean free path:  $L > \ell_e$ . Impurity scattering cannot be neglected and the electrons scatter in the sample undergoing a quantum diffusive dynamics. In a 2DEG at low temperature, quantum corrections to the conductance are mostly due to interference effects between two electrons scattered by the same impurity. These propagate along the same closed trajectory but in opposite directions — usually known as Cooperon correction. This process has its more spectacular manifestation in the so called WL (see Ref. [Akermans and Montambaux, 2007; Altshuler *et al.*, 1980; and Kramer, B. and MacKinnon, A., 1993]): it increases the effective scattering cross-section and therefore leads to a suppression of the conductivity. In the presence of a magnetic field  $\mathbf{B}$  perpendicular to the electron plane, the two waves acquire a phase difference  $\Delta\varphi = 2\frac{\phi_B}{\phi_0}$ , where  $\phi_B$  is the magnetic flux through the area enclosed by the electron trajectory, therefore quantum interference effects are suppressed and the conductance increases to the classical limit. Such a modification in conductivity, as a function of the magnetic field, is known as positive magneto conductivity (sometimes called negative magnetoresistance). This quantum correction to the resistivity can be explicitly calculated evaluating the Cooperon diagram. For a 2DEG in perpendicular magnetic field

$$\Delta\sigma(B) \propto \int_0^\infty dt Z_c(t, B) \left[ e^{-\frac{t}{\tau_\phi}} - e^{-\frac{t}{\tau_e}} \right], \quad (63)$$

where  $Z_c(t) = \frac{\phi_B}{\phi_0} \sinh^{-1} \left( \frac{4\pi DtB}{\phi_0} \right)$ , where  $\tau_\phi$  and  $\tau_e$  are phase coherence time and electron relaxation time and  $D$

is the 2D diffusion coefficient, respectively. In the limit of very weak magnetic field ( $B \ll \frac{\phi_0}{8\pi L_\phi^2}$ ) the magnetoresistance results to be:

$$\Delta R(B) \propto -\frac{\Delta\sigma(B)}{\sigma_0^2}, \quad (64)$$

thus negative, and the resistivity correction can be shown to be

$$\Delta\sigma(B) - \Delta\sigma(0) \propto \left(\frac{B}{B_\phi}\right)^2, \quad (65)$$

where  $B_\phi = \frac{\phi_0}{8\pi L_\phi^2}$  is a characteristic magnetic field corresponding to a flux quantum through an area of the order of  $\pi L_\phi^2$ . Measuring the negative magneto-resistance by applying a weak magnetic field is a very elegant way to probe weak localization correction to the conductance.

In the presence of SOI the WL correction changes its sign and the magnetoresistance results to be positive. Indeed, Cooperon correction is multiplied by a factor

$$\langle Q_{\text{SOI}}(t) \rangle = \frac{1}{2} \left[ 3e^{-\frac{t}{\tau_{\text{SOI}}}} - 1 \right],$$

where  $\tau_{\text{SOI}}$  sets the time scale of the SOI. For strong SOI, the exponential factor can be neglected at any time  $t$ , thus the factor  $\langle Q_{\text{SOI}} \rangle$  changes its sign and tends to  $-\frac{1}{2}$ . It results in a change of the sign of conventional weak localization correction. This phenomenon, theorised in Ref. (Hikami *et al.*, 1980), is called WAL. It was first observed in metal films in the presence of spin-orbit impurities (Bergman, 1984, 1982) and has been intensively studied in the recent past, both in 2DEGs (Miller *et al.*, 2003; Knap *et al.*, 1996; Koga *et al.*, 2002) and in large quantum dots (Zumbühl *et al.*, 2002; Aleiner and Fal'ko, 2001) made out of III-V semiconductors in the presence of RSOI [c.f. for instance figure 1(b)]. Recently, weak WAL measurements have been used to quantify the RSOI strength in oxide interfaces (Caviglia *et al.*, 2010) and TIs (Bardarson and Moore, 2013).

### C. Quantum wires with RSOI, superconductivity and magnetic field: The quest for Majorana Fermions

In his seminal article on the symmetric theory of electron and positron (Majorana, 1937), Ettore Majorana predicted the existence of selfadjoint fermions, *i.e.* half integer spin particles being their own antiparticle, as real solutions of Dirac equation. Since the first Majorana's prediction, the quest for Majorana Fermions (MFs) as elementary particles, in the high energy physics, is still open. Experiments on the neutrinoless double beta decay should unveil whether neutrinos are MFs or not, but to the date there is no answer to this question.

On the contrary, electrons in solid, under particular circumstances could recombine with holes — their relative anti-particles in the solid state language — in order to form self-adjoint excitations, *i.e.* Majorana quasiparticles.

Thus MFs (Kitaev, 2001, 2006) can appear as quasi-particle excitations in solid state systems. For instance, they are expected to show up as boundary states of the Kitaev model: a toy-model Hamiltonian describing 1D spineless  $p$ -wave superconductors.

Despite the fact that “solid state” MFs are not elementary particles, there is strong excitement in the condensed matter community, and many efforts are being devoted to their theoretical and experimental understanding. Of course, finding a novel elementary excitation in solids would have a fundamental significance on its own, however, in the case of MF there is strong interest also for its possible applications. Indeed MFs, due to their non-Abelian statistics (Read and Green, 2000; Ivanov, 2001; Stern *et al.*, 2004; Alicea *et al.*, 2011) can be considered as building blocks for topological quantum computation and for other alternative decoherence free quantum computational schemes (Nayak *et al.*, 2008).

Several mechanisms and devices have been proposed to isolate and detect MFs in meso/nano scaled devices (c.f. Refs. [Alicea, 2012 and Beenakker, 2013] for exhaustive reviews on the topic). However, to the date, while many efforts are focused on hybrid TI superconductor heterostructures (Veldhorst *et al.*, 2012; Galletti *et al.*, 2014a,b), (one of) the most promising device resorts on the use of III-V semiconducting nanowires with strong RSOI, in the proximity with conventional superconductors and in the presence of a magnetic field along the wire. This mechanism, early proposed in Refs. (Oreg *et al.*, 2010; Lutchyn *et al.*, 2010) has been later experimentally explored in Refs. [Das *et al.*, 2012; Mourik *et al.*, 2012; and Churchill *et al.*, 2013]. While strong hints of MF physics have been highlighted in Ref. [Mourik *et al.*, 2012], other recent articles [Churchill *et al.*, 2013; Linder *et al.*, 2010; Lee *et al.*, 2012; Liu *et al.*, 2012; Kells *et al.*, 2012; and Sun and Shah, 2015] offer alternative interpretation of the observed zero bias anomaly in terms of Kondo physics. Therefore an unambiguous “smoking-gun” experimental proof of the presence of MFs still has to come.

In the following, we will describe the theoretical background supporting the physics of MFs in quasi-1D nano wires with RSOI, magnetic field and proximity induced superconductivity. After that, we briefly review a recent experimental result.

For sake of simplicity we assume a single active channel in the nanowire and set  $\hbar = 1$ . The nanowire is along the  $x$  direction, therefore the transverse dynamics, along  $y$  direction, can be neglected. The Hamiltonian (15) describing the wire can be simplified as follows:

$$\mathcal{H}_{\text{MF}} = \sum_{\sigma\sigma'} \int_{-\infty}^{\infty} \psi_{\sigma}^{\dagger}(x) \left( \frac{p_x^2}{2m^*} - \alpha s_y p_x + g\mu_B B s_x - \mu \right) \psi_{\sigma'}(x) + \left( \Delta \psi_{\uparrow}^{\dagger}(x) \psi_{\downarrow}^{\dagger}(x) + \text{h.c.} \right) dx \quad (66)$$

where we have also added a Zeeman term due to the magnetic field  $B$  along the wire axis and the superconducting pairing potential  $\Delta$ . The operator  $\psi_{\sigma}^{\dagger}(x)$  creates an electron of spin  $\sigma$  at the position  $x$ . One can notice that the Hamiltonian (66) is not that of a 1D spinless  $p$ -wave superconductor, indeed we can easily recognise a local  $s$ -wave superconducting pairing and spinful electrons. However, by analysing the action of the various interaction terms in (66), we can show how to obtain an optimal choice of the parameter regime that would map (66) into the effective Hamiltonian of a 1D spinless  $p$ -wave superconductor.

We choose  $\Psi(x) = [u_{\uparrow}(x), u_{\downarrow}(x), v_{\downarrow}(x), -v_{\uparrow}(x)]$  as a Nambu spinor — where  $u/v$  are quasi electron/hole wave functions of momentum  $k$  (De Gennes, 1999). Thus the Bogolubov–de Gennes (BdG) Hamiltonian (66) in the  $\Psi(x)$  basis, can be recast in the simple matrix form:

$$\mathcal{H}_{\text{BdG}}(k) = \left( \frac{k^2}{2m^*} + i\alpha k s_y - \mu \right) \tau_z + E_z s_x + \Delta \tau_x \quad (67)$$

where the  $\tau_i$  Pauli matrices act in the Nambu space, while  $s_i$  act in the spin space and we have introduced the Zeeman energy  $E_z$ . In the absence of magnetic field and superconductivity, the conventional spin-degenerate parabolic band dispersion splits because of RSOI, into two parabolas crossing at  $k = 0$  [c.f. figure 5(e)]. The Zeeman coupling, splits the two bands at  $k = 0$  giving rise to the two bands  $E_+(k)$  and  $E_-(k)$  in figure 20(a). If the Fermi energy lies between the two bands, and the temperature is low enough that the higher band cannot be thermally populated, the system is effectively a 1D spinless system. The spin degree-of-freedom has been effectively quenched by the simultaneous action of the RSOI and Zeeman splitting. Only a single pseudo spin degree-of-freedom “ $-$ ” is involved here. In addition we have the  $s$ -wave superconducting pairing. Projecting the full Hamiltonian onto the  $E_-(k)$  band we can write down an effective Hamiltonian

$$\mathcal{H}_{\text{P}} = \sum_k \left( E_-(k) c_k^{\dagger} c_k + \Delta_-(k) c_k^{\dagger} c_k^{\dagger} + \text{h.c.} \right) \quad (68)$$

where  $\Delta_-(k) = i\alpha k_x \Delta (\alpha^2 k_x^2 + E_z^2)^{-1/2}$  is an effective superconducting pairing with the desired  $p$ -wave symmetry. Thus, the present problem is isomorphic to the Majorana wire considered by Kitaev [Kitaev, 2001]. That guarantees the presence of MFs as boundary excitations. However in the spirit of a review article, here we will not provide an explicit proof of the presence of MFs as boundary excitations of the wire, as it can be inferred by the isomorphism to Kitaev model and can also be found in Refs. [Oreg *et al.*, 2010 and Lutchyn *et al.*, 2010] but we

will provide the reader with simple symmetry arguments to qualitatively justify this result.

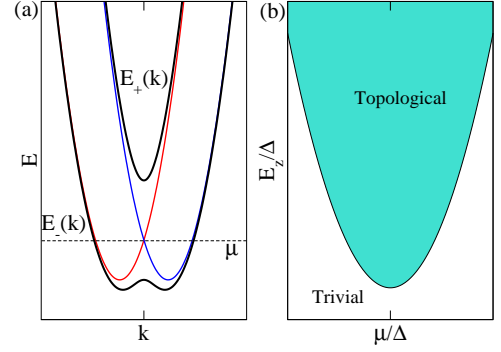


FIG. 20 (Colours online) (a) Free electrons band splitter by the simultaneous application of RSOI and Zeeman splitting. (b) Phase diagram of the model Hamiltonian (67).

Let us start by commenting on the spectrum. By squaring twice Hamiltonian (67) we obtain the dispersion relation for the two bands

$$E_{\pm}^2(k, \Delta) = E_z^2 + \Delta^2 + \xi_k^2 + (\alpha k)^2 \pm 2\sqrt{(E_z \Delta)^2 + (E_z \xi_k)^2 + (\alpha k \xi_k)^2}$$

where  $\xi_k = k^2/2m - \mu$ . The pairing  $\Delta$  plays two crucial roles. It opens a gap at the outer wings of the dispersion, where the Zeeman field is unimportant, and modifies the gap forming near  $k = 0$ . The former role eliminates the possibility of high-momentum gapless excitations, thus leaving only the chiral states near  $k = 0$  as low energy excitation. The latter role allows us to tune the topological phase transition essential for isolating MFs. One can easily verify that the Hamiltonian (67) satisfies particle-hole symmetry:

$$\Xi \mathcal{H}_{\text{BdG}}(k) \Xi^{-1} = \mathcal{H}_{\text{BdG}}(-k). \quad (69)$$

Hence, for each eigenstate of positive energy  $\mathcal{H}_{\text{BdG}}\psi = E\psi$  it exists a corresponding eigenstate  $\Xi\psi$  of opposite energy  $\mathcal{H}_{\text{BdG}}(\Xi\psi) = -E(\Xi\psi)$ . A linear combination  $\gamma = \psi + (\Xi\psi)$  is of course a selfadjoint Fermionic operator, *i.e.* a MF, however it is not an Hamiltonian eigenvector, unlike  $E = 0$  (Chamon *et al.*, 2010).

It can be shown that the presence of such zero energy selfadjoint eigenstate is indeed related to the sign of the gap at  $k = 0$

$$E_0 = |E_+(k, \Delta) - E_-(k, \Delta)|_{k=0} = |E_z - \sqrt{\Delta^2 + \mu^2}|$$



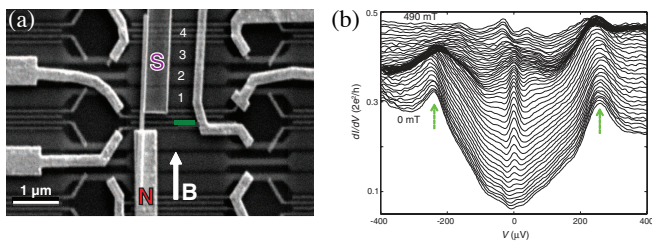


FIG. 21 (Colours online) (a): Experimental device. Electrons from a normal metal are injected in the InSb nanowire which is in contact with NbTiN superconductor. (b): Differential conductance traces as a function of the magnetic field. As soon as the topological criterion is satisfied, a zero bias peak appears. Courtesy of Ref. [Mourik *et al.*, 2012].

between the two excitation branches  $E_{\pm}(k, \Delta)$ . When the Zeeman energy  $E_z$  closes such gap, we have a quantum phase transition between two different physical regimes. At  $E_z > \sqrt{\Delta^2 + \mu^2}$  the gap is a Zeeman gap while in the opposite case the gap is due to superconductivity. Explicit construction of the zero energy excitation in the two cases shows that when the gap is magnetic, selfadjoint zero energy excitations, *i.e.* MFs can appear at the two boundaries of the wire. The phase diagram of such system is shown in figure 20(b). Out of the topological regime no MFs are expected, but the interesting physics of anomalous Josephson effect sets in (Yokoyama *et al.*, 2014; Campagnano *et al.*, 2014).

In Ref. [Mourik *et al.*, 2012] electrons are injected from a normal metal in an InSb nanowire in proximity with a NbTiN superconductor [see figure 21(a)] and the differential conductance is measured as a function of the magnetic field. As proposed in Ref. [Flensberg, 2010] electrons from the normal metal scatter against the zero energy MF thus revealing a zero bias anomaly in the differential conductance. In figure 21(b) we can see that as soon as the magnetic field is large enough to satisfy the topological criterion, a zero bias peak appears that is interpreted as the resonant scattering of electrons through the zero energy Majorana state. Experiments on these kind of structures are very challenging: on the one hand in order to satisfy the topological criterion, a sizable magnetic field is required, on the other hand, the same magnetic field can disrupt the very fragile superconductivity induced in the semiconducting nanowire by the proximity effect (Potter and Lee, 2011). Everything has to be finely tuned and the temperature has to be quite low, usually below 100mK. That is why many other proposals involving high critical temperature superconductors (Lucignano *et al.*, 2012, 2013) or completely different platforms (Nadj-Perge *et al.*, 2014; Hart *et al.*, 2014) have been investigated, too.

## VII. CONCLUSIONS AND OUTLOOK

In this review, we have inspected some of the most interesting theoretical and experimental results on spin dependent quantum transport in mesoscopic systems in the presence of RSOI. During the last decades a significant number of theories and experiments have appeared. The main goal of this review is to give an historical overview of the field so to address the reader directly to a more specialized literature. Rashba spin-orbit interaction turned out to be an invaluable tool to have access and to manipulate the electron spin degree of freedom, without breaking time reversal symmetry. From the applicative point of view, RSOI has been intensively studied to generate spin currents in semiconducting electron systems. Several mechanisms and devices for producing pure spin currents have been put forward. Among them, here we have focused our attention on quantum spin ratchet and spin pumping.

The physics associated to the RSOI can have very applicative oriented aspects but can also be of stimulus for addressing more fundamental issues of quantum mechanics, *e.g.*, quantum interference effects. These have been shown to be strongly affected by the RSOI, with the novelty of the phase factors of non-Abelian nature. We have seen how this additional phase can give rise to unexpected phenomena, such as the localisation in networks with nontrivial connectivity, or the appearance of anomalous peaks in the Fourier transform of the magneto conductance in quantum rings. Recently, new and exciting perspectives open, because RSOI is of fundamental relevance for the physics of topological insulators and of Majorana fermions in hybrid superconducting-semiconducting heterostructures that are currently attracting a lot of interest not only in the condensed matter community.

Besides charge and spin transport, SOI may have a relevant role also in heat transport which is the subject of *spin caloritronics* (Bauer *et al.*, 2012). Here, the combination of magnetic and non magnetic hybrid structure can lead to anomalous behaviour in the Seebeck and/or the Peltier coefficients (Gravier *et al.*, 2006; Flipse *et al.*, 2012). Deviation from standard behavior has been observed also in experiments on SLG (Zuev *et al.*, 2009; Wei *et al.*, 2009). Furthermore, it is well known that the most efficient media for creating thermoelectric devices are based on materials with a very strong SOI as HgTe, PbTe, Bi<sub>2</sub>Te<sub>3</sub>, and Bi/Sb alloys, Ref. (Delves, 1965; Biswas *et al.*, 2012). However, a complete understanding of the role of RSOI on these properties is still under investigation (Alomar and Sánchez, 2014; Rameshti and Moghaddam, 2015).

We are sure that in the future this field of research will be still thriving. Most probably new electronic and spintronic devices will come to the market with functionality that are associated to the physics of the RSOI. On

the other side, as it happened for the case of the Majorana quasiparticle research, RSOI could pave the way to fundamental research in order to investigate exotic phenomena that could get accessed by *quantum simulators* realised in solid state systems (Feynman, 1982).

## ACKNOWLEDGMENTS

We gratefully acknowledge all our collaborators on the subject of quantum transport in systems with Rashba spin-orbit interaction. In particular F.S. Bergeret, D. Gosálbez Martínez, V. Marigliano Ramaglia and A. Tagliacozzo that have carefully read the manuscript. D.B. acknowledges financial support from Ikerbasque Foundation of Science and the Alexander von Humboldt Foundation. P.L. acknowledges financial support from the FIRB project HybridNanoDev RBFR1236VV of Italian ministry of education.

## REFERENCES

- I. Žutić, J. Fabian, and S. das Sarma, *Rev. Mod. Phys.* **76**, 323 (2004).
- J. Sinova and I. Žutić, *Nat. Mater.* **11**, 368 (2012).
- M. N. Baibich, J. M. Broto, A. Fert, F. N. Van Dau, F. Petroff, P. Etienne, G. Creuzet, A. Friederich, and J. Chazelas, *Phys. Rev. Lett.* **61**, 2472 (1988).
- G. Binasch, P. Grünberg, F. Saurenbach, and W. Zinn, *Phys. Rev. B* **39**, 4828 (1989).
- B. Dieny, V. S. Speriosu, S. S. P. Parkin, B. A. Gurney, D. R. Wilhoit, and D. Mauri, *Phys. Rev. B* **43**, 1297 (1991).
- R. E. Camley and J. Barnaś, *Phys. Rev. Lett.* **63**, 664 (1989).
- J. S. Moodera, L. R. Kinder, T. M. Wong, and R. Meservey, *Phys. Rev. Lett.* **74**, 3273 (1995).
- M. Johnson and R. H. Silsbee, *Phys. Rev. B* **37**, 5326 (1988).
- S. Datta and B. Das, *Appl. Phys. Lett.* **56**, 665 (1990).
- E. Rashba, *Fiz. Tverd. Tela (Leningrad)* **2**, 1224 (1960), [*Sov. Phys. Solid State* **2**, 1109 (1960)].
- Yu A. Bychkov and E. Rashba, *J. Phys. C: Solid State Phys.* **17**, 6039 (1984).
- E. de Andrada e Silva, G. C. La Rocca, and F. Bassani, *Phys. Rev. B* **50**, 8523 (1994).
- E. de Andrada e Silva, G. C. La Rocca, and F. Bassani, *Phys. Rev. B* **55**, 16293 (1997).
- J. Nitta, T. Akazaki, H. Takayanagi, and T. Enoki, *Phys. Rev. Lett.* **78**, 1335 (1997).
- T. Schäpers, G. Engels, J. Lange, T. Klocke, M. Hollfelder, and H. Lüth, *J. Appl. Phys.* **83**, 4324 (1998).
- D. Grundler, *Phys. Rev. Lett.* **84**, 6074 (2000).
- J. B. Miller, D. M. Zumbühl, C. M. Marcus, Y. B. Lyanda-Geller, D. Goldhaber-Gordon, K. Campman, and A. C. Gossard, *Phys. Rev. Lett.* **90**, 076807 (2003).
- L. Meier, G. Salis, I. Shorubalko, E. Gini, S. Schoen, and K. Ensslin, *Nature Phys.* **3**, 650 (2007).
- G. Schmidt, D. Ferrand, L. W. Molenkamp, A. T. Filip, and B. J. van Wees, *Phys. Rev. B* **62**, R4790 (2000).
- H. Ohno, *Science* **281**, 951 (1998).
- H. C. Koo, J. H. Kwon, J. Eom, J. Chang, S. H. Han, and M. Johnson, *Science* **325**, 1515 (2009).
- P. Chuang, S.-C. Ho, L. W. Smith, F. Sfigakis, M. Pepper, C.-H. Chen, J.-C. Fan, J. P. Griffiths, I. Farrer, H. E. Beere, G. A. C. Jones, D. A. Ritchie, and T.-M. Chen, *Nat. Nanotechnol.* **10**, 35 (2014).
- Y. Aharonov and A. Casher, *Phys. Rev. Lett.* **53**, 319 (1984).
- J.-B. Yau, E. P. De Poortere, and M. Shayegan, *Phys. Rev. Lett.* **88**, 146801 (2002).
- J. Nitta, F. E. Meijer, and H. Takayanagi, *Appl. Phys. Lett.* **75**, 695 (1999).
- D. Frustaglia and K. Richter, *Phys. Rev. B* **69**, 235310 (2004).
- D. Bercioux, D. Frustaglia, and M. Governale, *Phys. Rev. B* **72**, 113310 (2005a).
- R. Capozza, D. Giuliano, P. Lucignano, and A. Tagliacozzo, *Phys. Rev. Lett.* **95**, 226803 (2005).
- P. Lucignano, D. Giuliano, and A. Tagliacozzo, *Phys. Rev. B* **76**, 045324 (2007).
- Y. Aharonov and D. Bohm, *Phys. Rev.* **115**, 485 (1959).
- K. Richter, *Physics* **5**, 22 (2012).
- M. D'yakonov and V. Perel', *Phys. Lett. A* **35**, 459 (1971).
- J. E. Hirsch, *Phys. Rev. Lett.* **83**, 1834 (1999).
- S. Murakami, N. Nagaosa, and S.-C. Zhang, *Science* **301**, 1348 (2003).
- J. Sinova, D. Culcer, Q. Niu, N. A. Sinitsyn, T. Jungwirth, and A. H. MacDonald, *Phys. Rev. Lett.* **92**, 126603 (2004).
- Y. K. Kato, R. C. Myers, A. C. Gossard, and D. D. Awschalom, *Science* **306**, 1910 (2004).
- V. Sih, R. C. Myers, Y. K. Kato, W. H. Lau, A. C. Gossard, and D. D. Awschalom, *Nature Phys.* **1**, 31 (2005a).
- J. Wunderlich, B. Kaestner, J. Sinova, and T. Jungwirth, *Phys. Rev. Lett.* **94**, 047204 (2005a).
- J. Wunderlich, B. Kaestner, J. Sinova, and T. Jungwirth, *Phys. Rev. Lett.* **94**, 047204 (2005b).
- V. Sih, W. H. Lau, R. C. Myers, V. R. Horowitz, A. C. Gossard, and D. D. Awschalom, *Phys. Rev. Lett.* **97**, 096605 (2006a).
- S. O. Valenzuela and M. Tinkham, *Nature* **442**, 176 (2006).
- B. H. Bransden and C. J. Joachain, *Physics of Atoms and Molecules* (Prentice Hall, 2003).
- J. J. Sakurai and J. J. Napolitano, *Modern Quantum Mechanics* (Pearson Higher Ed, 2014).
- G. Bastard, *Wave mechanics applied to semiconductor heterostructures* (Wiley-Interscience, 1988).
- K. Seeger, *Semiconductor Physics, An Introduction* (Springer, 2004).
- K. Samokhin, *Ann. Phys. (N.Y.)* **324** (2009), 10.1016/j.aop.2009.08.008.
- G. Dresselhaus, *Phys. Rev.* **100**, 580 (1955).
- R. Winkler, *Spin-orbit effects in two dimensional electron and hole systems* (Springer-Verlag Berlin, 2003).
- P. Lucignano, R. Raimondi, and A. Tagliacozzo, *Phys. Rev. B* **78**, 035336 (2008).
- D. Stein, K. v. Klitzing, and G. Weimann, *Phys. Rev. Lett.* **51**, 130 (1983).
- H. L. Stormer, Z. Schlesinger, A. Chang, D. C. Tsui, A. C. Gossard, and W. Wiegmann, *Phys. Rev. Lett.* **51**, 126 (1983).
- E. Akkermans and G. Montambaux, *Mesoscopic Physics of Electrons and Photons* (Cambridge Univ. Press, Cambridge, 2007).
- T. Schäpers, J. Knobbe, and V. Guzenko, *Phys. Rev. B* **69**, 235323 (2004).
- T. Schäpers, V. A. Guzenko, M. G. Pala, U. Zülicke, M. Governale, J. Knobbe, and H. Hardtdegen, *Phys. Rev. B* **74**, 81301 (2006).

- H. A. Nilsson, P. Caroff, C. Thelander, M. Larsson, J. B. Wagner, L.-E. Wernersson, L. Samuelson, and H. Q. Xu, *Nano Lett.* **9**, 3151 (2009).
- S. Nadj-Perge, V. S. Pribiag, J. W. G. van den Berg, K. Zuo, S. R. Plissard, E. P. A. M. Bakkers, S. M. Frolov, and L. P. Kouwenhoven, *Phys. Rev. Lett.* **108**, 166801 (2012).
- S. Estévez Hernández, M. Akabori, K. Sladek, C. Volk, S. Alagha, H. Hardtdegen, M. G. Pala, N. Demarina, D. Grützmacher, and T. Schäpers, *Phys. Rev. B* **82**, 235303 (2010).
- Y. Oreg, G. Refael, and F. von Oppen, *Phys. Rev. Lett.* **105**, 177002 (2010).
- R. M. Lutchyn, J. D. Sau, and S. das Sarma, *Phys. Rev. Lett.* **105**, 077001 (2010).
- A. Varykhalov, J. Sánchez-Barriga, A. Shikin, C. Biswas, E. Vescovo, A. Rybkin, D. Marchenko, and O. Rader, *Phys. Rev. Lett.* **101**, 157601 (2008).
- C. L. Kane and E. J. Mele, *Phys. Rev. Lett.* **95**, 146802 (2005a).
- C. L. Kane and E. Mele, *Phys. Rev. Lett.* **95**, 226801 (2005b).
- D. Huertas-Hernando, F. Guinea, and A. Brataas, *Phys. Rev. B* **74**, 155426 (2006).
- Y. S. Dedkov, M. Fonin, U. Rüdiger, and C. Laubschat, *Phys. Rev. Lett.* **100**, 107602 (2008).
- Z. Y. Li, Z. Q. Yang, S. Qiao, J. Hu, and R. Q. Wu, *Journal of Physics: Condensed Matter* **23**, 225502 (2011).
- D. Marchenko, A. Varykhalov, M. R. Scholz, G. Bihlmayer, E. I. Rashba, A. Rybkin, A. M. Shikin, and O. Rader, *Nat. Commun.* **3**, 1232 (2012).
- A. H. Castro Neto and F. Guinea, *Phys. Rev. Lett.* **103**, 026804 (2009).
- J. Klinovaja and D. Loss, *Phys. Rev. X* **3**, 011008 (2013).
- A. H. C. Neto, F. Guinea, N. M. R. Peres, K. S. Novoselov, and A. K. Geim, *Rev. Mod. Phys.* **81**, 109 (2009).
- R. v. Gelderen and C. M. Smith, *Phys. Rev. B* **81**, 125435 (2010).
- F. Guinea, *New J. Phys.* **12**, 083063 (2010).
- S. Konschuh, M. Gmitra, D. Kochan, and J. Fabian, *Phys. Rev. B* **85**, 115423 (2012).
- F. Mireles and J. Schliemann, *New J. Phys.* **14** (2012), 10.1088/1367-2630/14/9/093026.
- A. D. Martino, R. Egger, K. Hallberg, and C. Balseiro, *Phys. Rev. Lett.* **88**, 206402 (2002).
- A. D. Martino and R. Egger, *J. Phys. Condens. Matter* **17**, 5523 (2005).
- W. Izumida, K. Sato, and R. Saito, *J. Phys. Soc. Jpn.* **78**, 074707 (2009).
- F. Kuemmeth, S. Ilani, D. C. Ralph, and P. L. Mceuen, *Nature* **452**, 448 (2008).
- T. S. Jespersen, K. Grove-Rasmussen, J. Paaske, K. Muraki, T. Fujisawa, J. Nygard, and K. Flensberg, *Nature Phys.* **7**, 348 (2011).
- M. Z. Hasan and C. L. Kane, *Rev. Mod. Phys.* **82**, 3045 (2010).
- X.-L. Qi and S.-C. Zhang, *Rev. Mod. Phys.* **83**, 1057 (2011).
- Y. Ando, *J. Phys. Soc. Jpn.* **82**, 102001 (2013).
- B. A. Bernevig, T. L. Hughes, and S.-C. Zhang, *Science* **314**, 1757 (2006).
- B. A. Bernevig and S.-C. Zhang, *Phys. Rev. Lett.* **96**, 106802 (2006).
- M. König, A. Tschetschetkin, E. Hankiewicz, J. Sinova, V. Hock, V. Daumer, M. Schäfer, C. Becker, H. Buhmann, and L. Molenkamp, *Phys. Rev. Lett.* **96**, 076804 (2006).
- C. Liu, T. Hughes, X.-L. Qi, K. Wang, and S.-C. Zhang, *Phys. Rev. Lett.* **100**, 236601 (2008).
- I. Knez, R.-R. Du, and G. Sullivan, *Phys. Rev. Lett.* **107**, 136603 (2011).
- L. Fu, C. L. Kane, and E. Mele, *Phys. Rev. Lett.* **98**, 106803 (2007).
- L. Fu and C. L. Kane, *Phys. Rev. B* **76**, 045302 (2007).
- J. H. Bardarson and J. E. Moore, *Rep. Prog. Phys.* **76**, 056501 (2013).
- F. Qu, F. Yang, J. Chen, J. Shen, Y. Ding, J. Lu, Y. Song, H. Yang, G. Liu, J. Fan, Y. Li, Z. Ji, C. Yang, and L. Lu, *Phys. Rev. Lett.* **107**, 016802 (2011).
- A. Altland and S. B. D., *Altland Alexander, Simons Ben D. Condensed Matter Field Theory, Second Edition 2010* (Cambridge University Press, 2010).
- Y. V. Nazarov and Y. M. Blanter, *Quantum Transport Introduction to Nanoscience* (Cambridge University Press, 2013).
- H. B. Nielsen and M. Ninomiya, *Phys. Lett. B* **105**, 219 (1981).
- X. Wan, A. M. Turner, A. Vishwanath, and S. Y. Savrasov, *Phys. Rev. B* **83**, 205101 (2011).
- S. Murakami, *New J. Phys.* **9**, 356 (2007).
- B.-J. Yang and Y. B. Kim, *Phys. Rev. B* **82**, 085111 (2010).
- O. Vafek and A. Vishwanath, *Annu. Rev. Condens. Matter Phys.* **5**, 83 (2014).
- P. Hosur and X. Qi, *C. R. Acad. Sci.* **14**, 857 (2013).
- A. V. Moroz and C. H. W. Barnes, *Phys. Rev. B* **60**, 14272 (1999).
- M. Governale and U. Zülicke, *Phys. Rev. B* **66**, 073311 (2002).
- C. Perroni, D. Bercioux, V. Ramaglia, and V. Cataudella, *J. Phys.: Condens. Matter* **19**, 186227 (2007).
- S. Smirnov, D. Bercioux, and M. Grifoni, *EPL* **80**, 27003 (2007).
- D. Sánchez and L. Serra, *Phys. Rev. B* **74**, 153313 (2006).
- Eto Mikio, Hayashi Tetsuya, and Kurotani Yuji, *J. Phys. Soc. Jpn.* **74**, 1934 (2005).
- M. Zarea and N. Sandler, *Phys. Rev. B* **79**, 165442 (2009).
- P. Rakyta, A. Kormányos, and J. Cserti, *Phys. Rev. B* **82**, 113405 (2010).
- L. Lenz, D. F. Urban, and D. Bercioux, *Eur. Phys. J. B* **86**, 502 (2013).
- D. Bercioux and A. De Martino, *Phys. Rev. B* **81**, 165410 (2010).
- M. Born and E. Wolf, *Principles of Optics: Electromagnetic Theory of Propagation, Interference and Diffraction of Light* (Cambridge University Press, 1999).
- V. Ramaglia, D. Bercioux, V. Cataudella, G. De Filippis, C. Perroni, and F. Ventriglia, *Eur. Phys. J. B* **36**, 365 (2003).
- M. Khodas, A. Shekhter, and A. M. Finkel'stein, *Phys. Rev. Lett.* **92**, 086602 (2004).
- V. Ramaglia, D. Bercioux, V. Cataudella, G. De Filippis, and C. Perroni, *J. Phys.: Condens. Matter* **16**, 9143 (2004).
- A. Messiah, *Quantum Mechanics* (Dover Publ Inc, 2014).
- G. E. Blonder, M. Tinkham, and T. M. Klapwijk, *Phys. Rev. B* **25**, 4515 (1982).
- i. d. I. Adagideli, V. Lutsker, M. Scheid, P. Jacquod, and K. Richter, *Phys. Rev. Lett.* **108**, 236601 (2012).
- C. Gorini, R. Raimondi, and P. Schwab, *Phys. Rev. Lett.* **109**, 246604 (2012).
- L. Lenz and D. Bercioux, *EPL* **96**, 27006 (2011).
- A. T. Costa, M. S. Ferreira, T. Hallam, G. S. Duesberg, and A. H. C. Neto, *EPL* **104**, 47001 (2013).
- M. Scheid, D. Bercioux, and K. Richter, *New J. Phys* **9**, 401

- (2007a).
- E. I. Rashba, *Phys. E* **34**, 31 (2006).
- E. I. Rashba, *Phys. Rev. B* **68**, 241315 (2003).
- B. Nikolić, L. Žárbo, and S. Souma, *Phys. Rev. B* **73**, 075303 (2006).
- A. Kiselev and K. Kim, *Phys. Rev. B* **71**, 153315 (2005).
- E. M. Hankiewicz, L. W. Molenkamp, T. Jungwirth, and J. Sinova, *Phys. Rev. B* **70**, 241301 (2004).
- W. Ren, Z. Qiao, J. Wang, Q. Sun, and H. Guo, *Phys. Rev. Lett.* **97**, 066603 (2006).
- J. H. Bardarson, i. d. I. Adagideli, and P. Jacquod, *Phys. Rev. Lett.* **98**, 196601 (2007).
- C. Brüne, A. Roth, H. Buhmann, E. M. Hankiewicz, L. W. Molenkamp, J. Maciejko, X.-L. Qi, and S.-C. Zhang, *Nature Phys.* **8**, 485 (2012).
- Q.-f. Sun and X. C. Xie, *Phys. Rev. B* **72**, 245305 (2005).
- J. Shi, P. Zhang, D. Xiao, and Q. Niu, *Phys. Rev. Lett.* **96**, 076604 (2006).
- Z. An, F. Q. Liu, Y. Lin, and C. Liu, *Sci. Rep.* **2**, 388 (2012).
- P. Reimann, *Phys. Rep.* **361**, 57 (2002).
- P. Hänggi and F. Marchesoni, *Rev. Mod. Phys.* **81**, 387 (2009).
- P. Reimann, M. Grifoni, and P. Hänggi, *Phys. Rev. Lett.* **79**, 10 (1997).
- H. Linke, T. Humphrey, A. Löfgren, A. Sushkov, R. Newbury, R. Taylor, and P. Omling, *Science* **286**, 2314 (1999).
- H. Linke, T. R. Humphrey, P. R. Lindelof, A. Löfgren, R. Newbury, P. Omling, A. O. Sushkov, R. P. Taylor, and H. Xu, *Appl. Phys. A* **75**, 2037 (2002).
- M. Scheid, A. Pfund, D. Bercioux, and K. Richter, *Phys. Rev. B* **76**, 195303 (2007b).
- M. Scheid, D. Bercioux, and K. Richter, *Chem. Phys.* **375**, 276 (2010).
- Y. S. Ang, Z. Ma, and C. Zhang, *Sci. Rep.* **5**, 7872 (2015).
- S. Smirnov, D. Bercioux, M. Grifoni, and K. Richter, *Phys. Rev. Lett.* **100**, 230601 (2008a).
- S. Smirnov, D. Bercioux, M. Grifoni, and K. Richter, *Phys. Rev. B* **78**, 245323 (2008b).
- S. Smirnov, D. Bercioux, M. G., and K. Richter, *Phys. Rev. B* **80**, 201310 (2009).
- M. Scheid, M. Wimmer, D. Bercioux, and K. Richter, *Phys. Status Solidi (c)* **3**, 4235 (2006).
- M. V. Costache and S. O. Valenzuela, *Science* **330**, 1645 (2010).
- P. W. Brouwer, *Phys. Rev. B* **58**, R10135 (1998).
- M. Governale, F. Taddei, and R. Fazio, *Phys. Rev. B* **68**, 155324 (2003).
- E. Mucciolo, C. Chamon, and C. Marcus, *Phys. Rev. Lett.* **89**, 146802 (2002).
- S. Watson, R. Potok, C. Marcus, and V. Umansky, *Phys. Rev. Lett.* **91**, 258301 (2003).
- D. Bercioux, D. F. Urban, F. Romeo, and R. Citro, *Appl. Phys. Lett.* **101**, 122405 (2012).
- R. Citro, F. Romeo, and N. Andrei, *Phys. Rev. B* **84**, 161301 (2011).
- A. Inhofer and D. Bercioux, *Phys. Rev. B* **88**, 235412 (2013).
- P. P. Hofer and M. Büttiker, *Phys. Rev. B* **88**, 241308 (2013).
- P. P. Hofer, H. Aramberri, C. Schenke, and P. A. L. Delplace, *EPL* **107**, 27003 (2014).
- F. Mireles and G. Kirczenow, *Phys. Rev. B* **64**, 024426 (2001).
- C. W. Groth, M. Wimmer, A. R. Akhmerov, and X. Waintal, *New J. Phys.* **16**, 063065 (2014).
- J.-S. Jeong and H.-W. Lee, *Phys. Rev. B* **74**, 195311 (2006).
- M. G. Pala, M. Governale, J. K. onig, U. Z. ulicke, and G. Iannaccone, *Phys. Rev. B* **69**, 045304 (2004).
- F. J. Jedema, A. T. Filip, and B. J. van Wees, *Nature* **410**, 345 (2001).
- F. J. Jedema, H. B. Heersche, A. T. Filip, J. J. A. Baselmans, and B. J. van Wees, *Nature* **416**, 713 (2002).
- X. Lou, C. Adelman, S. A. Crooker, E. S. Garlid, J. Zhang, K. S. M. Reddy, S. D. Flexner, C. J. Palmstrøm, and P. A. Crowell, *Nature Phys.* **3**, 197 (2007).
- T. Bergsten, T. Kobayashi, Y. Sekine, and J. Nitta, *Phys. Rev. Lett.* **97**, 196803 (2006).
- F. Nagasawa, J. Takagi, Y. Kunihashi, M. Kohda, and J. Nitta, *Phys. Rev. Lett.* **108**, 086801 (2012).
- F. Nagasawa, D. Frustaglia, H. Saarikoski, K. Richter, and J. Nitta, *Nat. Commun.* **4**, 2526 (2013).
- B. Sutherland, *Phys. Rev. B* **34**, 5208 (1986).
- D. Bercioux, M. Governale, V. Cataudella, and V. Ramaglia, *Phys. Rev. B* **72**, 075305 (2005b).
- D. Bercioux, D. F. Urban, H. Grabert, and W. Haeusler, *Phys. Rev. A* **80**, 063603 (2009).
- D. Bercioux, N. Goldman, and D. F. Urban, *Phys. Rev. A* **83**, 023609 (2011).
- J. Vidal, R. Mosseri, and B. Doucot, *Phys. Rev. Lett.* **81**, 5888 (1998).
- J. Vidal, B. Doucot, R. Mosseri, and P. Butaud, *Phys. Rev. Lett.* **85**, 3906 (2000a).
- C. C. Naud, G. G. Faini, and D. D. Mailly, *Phys. Rev. Lett.* **86**, 5104 (2001).
- D. Bercioux, M. Governale, V. Cataudella, and V. M. Ramaglia, *Phys. Rev. Lett.* **93**, 056802 (2004).
- J. Vidal, G. Montambaux, and B. Doucot, *Phys. Rev. B* **62**, 16294 (2000b).
- C. C. Abilio, P. Butaud, T. Fournier, B. Pannetier, J. Vidal, S. Tedesco, and B. Dalzotto, *Phys. Rev. Lett.* **83**, 5102 (1999).
- N. Hatano, R. Shirasaki, and H. Nakamura, *Phys. Rev. A* **75**, 032107 (2007).
- A. Aharony, O. Entin-Wohlman, Y. Tokura, and S. Katsumoto, *Phys. Rev. B* **78**, 125328 (2008).
- A. Aharony, Y. Tokura, G. Z. Cohen, O. Entin-Wohlman, and S. Katsumoto, *Phys. Rev. B* **84**, 035323 (2011).
- S. Matityahu, A. Aharony, O. Entin-Wohlman, and S. Katsumoto, *Phys. Rev. B* **87**, 205438 (2013a).
- S. Matityahu, A. Aharony, O. Entin-Wohlman, and S. Tarucha, *New J. Phys.* **15**, 125017 (2013b).
- S. Washburn and R. A. Webb, *Rep. Prog. Phys.* **55**, 1311 (1992).
- J. Anandan, *Science* **297**, 1656 (2002).
- H. Saarikoski, J. E. Vázquez-Lozano, J. P. Baltanás, F. Nagasawa, J. Nitta, and D. Frustaglia, arXiv:1412.5487 (2014).
- M. V. Berry, *Proc. R. Soc. Lond. A* **392**, 45 (1984).
- Y. Aharonov and J. Anandan, *Phys. Rev. Lett.* **58**, 1593 (1987).
- J. Nitta, T. Koga, and F. E. Meijer, *Phys. E* **18**, 143 (2003).
- F. Meijer, J. Nitta, T. Koga, A. Morpurgo, and T. Klapwijk, *Phys. E* **22**, 402 (2004).
- A. F. Morpurgo, J. P. Heida, T. M. Klapwijk, B. J. van Wees, and G. Borghs, *Phys. Rev. Lett.* **80**, 1050 (1998).
- B. Habib, E. Tutuc, and M. Shayegan, *Appl. Phys. Lett.* **90**, 152104 (2007).
- D. Loss, P. Goldbart, and A. V. Balatsky, *Phys. Rev. Lett.* **65**, 1655 (1990).
- H.-A. Engel and D. Loss, *Phys. Rev. B* **62**, 10238 (2000).
- Y. Tserkovnyak and A. Brataas, *Phys. Rev. B* **76**, 155326 (2007).

- B. Molnár, F. M. Peeters, and P. Vasilopoulos, *Phys. Rev. B* **69**, 155335 (2004).
- R. Citro, F. Romeo, and M. Marinaro, *Phys. Rev. B* **74**, 115329 (2006).
- V. M. Ramaglia, V. Cataudella, G. De Filippis, and C. A. Perroni, *Phys. Rev. B* **73**, 155328 (2006).
- S. Souma and B. K. Nikolić, *Phys. Rev. B* **70**, 195346 (2004).
- B. H. Wu and J. C. Cao, *Phys. Rev. B* **74**, 115313 (2006).
- F. E. Meijer, A. F. Morpurgo, and T. M. Klapwijk, *Phys. Rev. B* **66**, 033107 (2002).
- T. Jungwirth, J. Wunderlich, and K. Olejnik, *Nat. Mater.* **11**, 382 (2012).
- G. Vignale, *J. Supercond. Nov. Magn.* **23**, 3 (2010).
- M. I. D'yakonov and V. I. Perel, *Pris'ma Zh. Eksp. Teor. Fiz.* **13**, 657 (1971a), [*Sov. Phys. JETP Lett.* **13**, 467 (1971)].
- M. I. D'yakonov and V. I. Perel, *Phys. Lett. A* **83**, 1834 (1971b).
- E. Hall, *Philos. Mag.* **12**, 157 (1881).
- R. Karplus and J. Luttinger, *Phys. Rev. B* **95**, 1154 (1954).
- P. Nozieres and C. Lewiner, *J. Phys. (Paris)* **34**, 901 (1973).
- N. Nagaosa, J. Sinova, S. Onoda, A. H. MacDonald, and N. P. Ong, *Rev. Mod. Phys.* **82**, 1539 (2010).
- S. Zhang, *Phys. Rev. Lett.* **85**, 393 (2000).
- A. Crépieux and P. Bruno, *Phys. Rev. B* **64**, 014416 (2001).
- R. Raimondi and P. Schwab, *Phys. Rev. B* **71**, 033311 (2005).
- O. V. Dimitrova, *Phys. Rev. B* **71**, 245327 (2005).
- T. Tanaka, H. Kontani, M. Naito, T. Naito, D. S. Hirashima, K. Yamada, and J. Inoue, *Phys. Rev. B* **77**, 165117 (2008).
- H.-A. Engel, E. I. Rashba, and B. I. Halperin, *Phys. Rev. Lett.* **98**, 036602 (2007).
- Q. Lin, S. Liu, and X. Lei, *Appl. Phys. Lett.* **88**, 122105 (2006).
- M. Milletari, R. Raimondi, and P. Schwab, *EPL* **82**, 67005 (2008).
- R. Raimondi, C. Gorini, P. Schwab, and M. Dzierzawa, *Phys. Rev. B* **74**, 035340 (2006).
- C. Gorini, P. Schwab, M. Dzierzawa, and R. Raimondi, *Phys. Rev. B* **78**, 125327 (2008).
- J. Wunderlich, B. Kaestner, J. Sinova, and T. Jungwirth, *Phys. Rev. Lett.* **94**, 047204 (2005c).
- S. Matsuzaka, Y. Ohno, and H. Ohno, *Phys. Rev. B* **80**, 241305 (2009).
- K. Nomura, J. Wunderlich, J. Sinova, B. Kaestner, A. H. MacDonald, and T. Jungwirth, *Phys. Rev. B* **72**, 245330 (2005).
- V. Sih, R. C. Myers, Y. K. Kato, W. H. Lau, A. C. Gossard, and D. D. Awschalom, *Nature Phys.* **1**, 31 (2005b).
- V. Sih, W. H. Lau, R. C. Myers, V. R. Horowitz, A. Gossard, and D. D. Awschalom, *Phys. Rev. Lett.* **97**, 096605 (2006b).
- N. P. Stern, S. Ghosh, G. Xiang, M. Zhu, N. Samarth, and D. D. Awschalom, *Phys. Rev. Lett.* **97**, 126603 (2006).
- N. P. Stern, D. W. Steuerma, S. Mack, A. C. Gossard, and D. D. Awschalom, *Appl. Phys. Lett.* **91**, 062109 (2007).
- N. P. Stern, D. W. Steuerma, S. Mack, A. C. Gossard, and D. D. Awschalom, *Nature Phys.* **4**, 843 (2008).
- H. Zhao, E. J. Loren, H. M. v. Driel, and A. L. Smirl, *Phys. Rev. Lett.* **96**, 246601 (2006).
- N. Okamoto, H. Kurebayashi, T. Trypiniotis, I. Farrer, D. A. Ritchie, E. Saitoh, J. Sinova, J. Masek, T. Jungwirth, and C. H. W. Barnes, *Nat. Mater.* **13**, 932 (2014).
- B. L. Altshuler, D. Khmel'nitzkii, A. I. Larkin, and P. A. Lee, *Phys. Rev. B* **22**, 5142 (1980).
- Kramer, B. and MacKinnon, A., *Rep. Prog. Phys.* **56**, 1469 (1993).
- S. Hikami, A. Larkin, and Y. Nagaoka, *Prog. Theor. Phys.* **63**, 707 (1980).
- G. Bergman, *Phys. Rep.* **107**, 1 (1984).
- G. Bergman, *Phys. Rev. Lett.* **48**, 1046 (1982).
- W. Knap, C. Skierbiszewski, A. Zduniak, E. Litwin-Staszewska, D. Bertho, F. Kobbi, J. L. Robert, G. E. Pikus, F. G. Pikus, S. V. Iordanskii, V. Mosser, K. Zekentes, and Y. B. Lyanda-Geller, *Phys. Rev. B* **53**, 3912 (1996).
- T. Koga, J. Nitta, T. Akazaki, and H. Takayanagi, *Phys. Rev. Lett.* **89**, 046801 (2002).
- D. M. Zumbühl, J. B. Miller, C. M. Marcus, K. Campman, and A. C. Gossard, *Phys. Rev. Lett.* **89**, 276803 (2002).
- I. L. Aleiner and V. I. Fal'ko, *Phys. Rev. Lett.* **87**, 256801 (2001).
- A. D. Caviglia, M. Gabay, S. Gariglio, N. Reyren, C. Cancellieri, and J.-M. Triscone, *Phys. Rev. Lett.* **104**, 126803 (2010).
- E. Majorana, *Nuovo Cimento* **14**, 171 (1937).
- A. Kitaev, *Phys. Usp.* **44**, 131 (2001).
- A. Kitaev, *Annals of Physics* **321**, 2 (2006).
- N. Read and D. Green, *Phys. Rev. B* **61**, 10267 (2000).
- D. A. Ivanov, *Phys. Rev. Lett.* **86**, 268 (2001).
- A. Stern, F. v. Oppen, and E. Mariani, *Phys. Rev. B* **70**, 205338 (2004).
- J. Alicea, Y. Oreg, G. Rafael, F. v. Oppen, and M. P. A. Fisher, *Nature Phys.* **7**, 412 (2011).
- C. Nayak, S. Simon, A. Stern, M. Freedman, and S. d. Sarma, *Rev. Mod. Phys.* **80**, 1083 (2008).
- J. Alicea, *Rep. Prog. Phys.* **75**, 076501 (2012).
- C. W. J. Beenakker, *Annu. Rev. Cond. Mat. Phys.* **4**, 112 (2013).
- M. Veldhorst, M. Snelder, M. Hoek, T. Gang, V. K. Guduru, X. L. Wang, U. Zeitler, W. G. van der Wiel, A. A. Golubov, H. Hilgenkamp, and A. Brinkman, *Nat. Mater.* **11**, 417 (2012).
- L. Galletti, S. Charpentier, M. Iavarone, P. Lucignano, D. Massarotti, R. Arpaia, Y. Suzuki, K. Kadowaki, T. Bauch, A. Tagliacozzo, F. Tafuri, and F. Lombardi, *Phys. Rev. B* **89**, 134512 (2014a).
- L. Galletti, S. Charpentier, P. Lucignano, D. Massarotti, R. Arpaia, F. Tafuri, T. Bauch, Y. Suzuki, A. Tagliacozzo, K. Kadowaki, and F. Lombardi, *Phys. C* **503**, 162 (2014b).
- A. Das, Y. Ronen, Y. Most, Y. Oreg, M. Heiblum, and H. Shtrikman, *Nature Phys.* **8**, 887 (2012).
- V. Mourik, K. Zuo, S. Frolov, S. Plissard, E. Bakkers, and L. Kouwenhoven, *Science* **336**, 1003 (2012).
- H. O. H. Churchill, V. Fatemi, K. Grove-Rasmussen, M. T. Deng, P. Caroff, H. Q. Xu, and C. M. Marcus, *Phys. Rev. B* **87**, 241401 (2013).
- J. Linder, Y. Tanaka, T. Yokoyama, A. Sudbø, and N. Nagaosa, *Phys. Rev. Lett.* **104**, 067001 (2010).
- E. J. H. Lee, X. Jiang, R. Aguado, G. Katsaros, C. M. Lieber, and S. D. Franceschi, *Phys. Rev. Lett.* **109**, 186802 (2012).
- J. Liu, A. C. Potter, K. T. Law, and P. A. Lee, *Phys. Rev. Lett.* **109**, 267002 (2012).
- G. Kells, D. Meidan, and P. W. Brouwer, *Phys. Rev. B* **86**, 100503 (2012).
- K. Sun and N. Shah, *Phys. Rev. B* **91**, 144508 (2015).
- P. G. De Gennes, *Superconductivity of metals and alloys*, Advanced Book Classics (Westview press, 1999).
- C. Chamon, R. Jackiw, Y. Nishida, S. Y. Pi, and L. Santos, *Phys. Rev. B* **81**, 224515 (2010).
- T. Yokoyama, M. Eto, and Y. V. Nazarov, *Phys. Rev. B* **89**, 195407 (2014).

- G. Campagnano, P. Lucignano, D. Giuliano, and A. Tagliacozzo, *J. Phys.: Condens. Matter* **27**, 205301 (2014).
- K. Flensberg, *Phys. Rev. B* **82**, 180516 (2010).
- A. C. Potter and P. A. Lee, *Phys. Rev. B* **83**, 184520 (2011).
- P. Lucignano, A. Mezzacapo, F. Tafuri, and A. Tagliacozzo, *Phys. Rev. B* **86**, 144513 (2012).
- P. Lucignano, F. Tafuri, and A. Tagliacozzo, *Phys. Rev. B* **88**, 184512 (2013).
- S. Nadj-Perge, I. K. Drozdov, J. Li, H. Chen, S. Jeon, J. Seo, A. H. MacDonald, B. A. Bernevig, and A. Yazdani, *Science* **346**, 602 (2014).
- S. Hart, H. Ren, T. Wagner, P. Leubner, M. Mühlbauer, C. Brüne, H. Buhmann, L. W. Molenkamp, and A. Yacoby, *Nature Phys.* **10**, 638 (2014).
- G. E. W. Bauer, E. Saitoh, and B. J. van Wees, *Nat. Mater.* **11** (2012), 10.1038/nmat3301.
- L. Gravier, S. Serrano-Guisan, F. Reuse, and J.-P. Ansermet, *Phys. Rev. B* **73**, 052410 (2006).
- J. Flipse, F. L. Bakker, A. Slachter, F. K. Dejene, and B. J. van Wees, *Nat. Nanotechnol.* **7**, 166 (2012).
- Y. M. Zuev, W. Chang, and P. Kim, *Phys. Rev. Lett.* **102**, 096807 (2009).
- P. Wei, W. Bao, Y. Pu, C. N. Lau, and J. Shi, *Phys. Rev. Lett.* **102**, 166808 (2009).
- R. T. Delves, *Reports On Progress In Physics* **28**, 249 (1965).
- K. Biswas, J. He, I. D. Blum, C.-I. Wu, T. P. Hogan, D. N. Seidman, V. P. Dravid, and M. G. Kanatzidis, *Nature* **489**, 414 (2012).
- M. I. Alomar and D. Sánchez, *Phys. Rev. B* **89**, 115422 (2014).
- B. Z. Rameshti and A. G. Moghaddam, *Phys. Rev. B* **91**, 155407 (2015).
- R. P. Feynman, *Int. J. Theor. Phys.* **21**, 467 (1982).