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A unified theory of spin-relaxation due to spin-orbit coupling in metals and semiconductors

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Spintronics is an emerging paradigm with the aim to replace conventional electronics by using electron spins as information carriers. Its utility relies on the magnitude of the spin-relaxation, which is dominated by spin-orbit coupling (SOC). Yet, SOC induced spin-relaxation in metals and semiconductors is discussed for the seemingly orthogonal cases when inversion symmetry is retained or broken by the so-called Elliott-Yafet and D'yakonov-Perel' spin-relaxation mechanisms, respectively. We unify the two theories on general grounds for a generic two-band system containing intra- and inter-band SOC. While the previously known limiting cases are recovered, we also identify parameter domains when a crossover occurs between them, i.e. when an inversion symmetry broken state evolves from a D'yakonov-Perel' to an Elliott-Yafet type of spin-relaxation and conversely for a state with inversional symmetry. This provides an ultimate link between the two mechanisms of spin-relaxation.

A future spintronics device would perform calculations and store information using the spin-degree of freedom of electrons with a vision to eventually replace conventional electronics^{1–3}. A spin-polarized ensemble of electrons whose spin-state is manipulated in a transistor-like configuration and is read out with a spin-detector (or *spin-valve*) would constitute an elemental building block of a spin-transistor. Clearly, the utility of spintronics relies on whether the spin-polarization of the electron ensemble can be maintained sufficiently long. The basic idea behind spintronics is that coherence of a spin-ensemble persists longer than the coherence of electron momentum due to the relatively weaker coupling of the spin to the environment. The coupling is relativistic and has thus a relatively weak effect known as spin-orbit coupling (SOC).

The time characterizing the decay of spin-polarization is the so-called spin-relaxation time (often also referred to as spin-lattice relaxation time), τ_s . It can be measured either using electron spin-resonance spectroscopy (ESR)⁴ or in spin-transport experiments^{5,6}. Much as the theory and experiments of spin-relaxation measurements are developed, it remains an intensively studied field for novel materials; e.g. the value of τ_s is the matter of intensive theoretical studies^{7–13} and spin-transport experiments^{14–17} in graphene at present.

The two most important spin-relaxation mechanisms in metals and semiconductors are the so-called Elliott-Yafet (EY) and the D'yakonov-Perel' (DP) mechanisms. These are conventionally discussed along disjoint avenues, due to reasons described below. Although the interplay between these mechanisms has been studied in semiconductors^{3,18–20}, no attempts have been made to unify their descriptions. We note that a number of other spin-relaxation mechanisms, e.g. that involving nuclear-hyperfine interaction, are known^{2,3}.

The EY theory^{21,22} describes spin-relaxation in metals and semiconductors with inversion symmetry. Therein, the SOC does not split the spin-up/down states ($|\uparrow\rangle, |\downarrow\rangle$) in the conduction band, however the presence of a near lying band weakly mixes these states while maintaining the energy degeneracy. The nominally up state reads: $|\hat{\uparrow}\rangle = a_k|\uparrow\rangle + b_k|\downarrow\rangle$ (here a_k, b_k are band structure dependent) and $b_k/a_k = L/\Delta$, where L is the SOC matrix element between the adjacent bands and Δ is their separation. E.g. in alkali metals $L/\Delta \approx 10^{-2}..10^{-3}$ (Ref. 22). Elliott showed using first order time-dependent perturbation theory that an electron can flip its spin with probability $(L/\Delta)^2$ at a momentum scattering event. As a result, the spin scattering rate ($\Gamma_s = \hbar/2\tau_s$) reads:

$$\Gamma_{s,EY} \approx \left(\frac{L}{\Delta}\right)^2 \Gamma, \quad (1)$$



where $\Gamma = \hbar/2\tau$ is the quasi-particle scattering rate with τ being the corresponding momentum scattering (or relaxation) time. This mechanism is schematically depicted in Fig. 1a.

For semiconductors with zinc-blende crystal structure, such as e.g. GaAs, the lack of inversion symmetry results in an efficient relaxation mechanism, the D'yakonov-Perel' spin-relaxation²³. Therein, the spin-up/down energy levels in the conduction bands are split. The splitting acts on the electrons as if an internal, k -dependent magnetic field would be present, around which the electron spins precess with a Larmor frequency of $\Omega(k) = \mathcal{L}(k)/\hbar$. Here $\mathcal{L}(k)$ is the energy scale for the inversion symmetry breaking induced SOC. Were no momentum scattering present, the electron energies would acquire a distribution according to $\Omega(k)\hbar$. In the presence of momentum scattering which satisfies $\Omega(k)\cdot\tau \ll 1$, the distribution is "motionally-narrowed" and the resulting spin-relaxation rate reads:

$$\Gamma_{s,DP} \approx \frac{\mathcal{L}^2}{\Gamma}. \quad (2)$$

This situation is depicted in Fig. 1b. Clearly, the EY and DP mechanisms result in different dependence on Γ which is often used for the empirical assignment of the relaxation mechanism²⁴.

The observation of an anomalous temperature dependence of the spin-relaxation time in MgB_2 ²⁵ and the alkali fullerenes²⁶ and the development of a generalization of the EY theory high-lighted that the spin-relaxation theory is not yet complete. In particular, the first order perturbation theory of Elliott breaks down when the quasi-particle scattering rate is not negligible compared to the other energy scales. One expects similar surprises for the DP theory when the magnitude of e.g. the Zeeman energy is considered in comparison to the other relevant energy scales.

Herein, we develop a general and robust theory of spin-relaxation in metals and semiconductors including SOC between different bands and the same bands, provided the crystal symmetry allows for the latter. We employ the Mori-Kawasaki theory which considers the propagation of the electrons under the perturbation of the SOC. We obtain a general result which contains both the EY and the DP mechanisms as limits when the quasi-particle scattering and the magnetic field are small. Interesting links are recognized between the two mechanisms when these conditions are violated: the EY mechanism appears to the DP-like when Γ is large compared to Δ and the DP mechanism appears to be EY-like when the Zeeman energy is larger than Γ . Qualitative explanations are provided for these analytically observed behaviors.

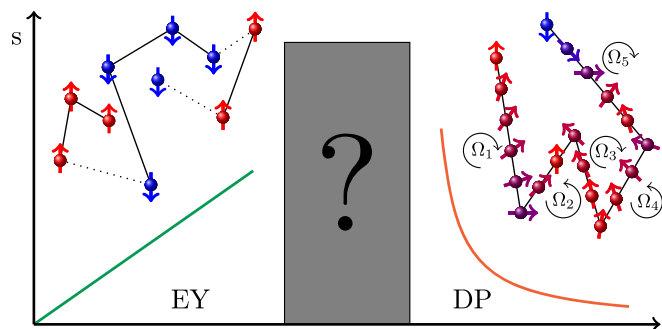


Figure 1 | Schematics of the Elliott-Yafet and the D'yakonov-Perel' mechanisms: $\Gamma_s \propto \Gamma$ in the EY scenario and spin-scattering occurs rarely (typically for every $10^4 \dots 10^6$ th momentum scattering in alkali metals), whereas the spin direction continuously precesses around the internal magnetic field due to SOC in the DP scenario, resulting in $\Gamma_s \propto 1/\Gamma$. It is the topic of the present paper, how these two distinct regimes are related to each other.

Results

The minimal model of spin-relaxation is a four-state (two bands with spin) model Hamiltonian for a two-dimensional electron gas (2DEG) in a magnetic field, which reads:

$$\mathcal{H} = \mathcal{H}_0 + \mathcal{H}_Z + \mathcal{H}_{\text{scatt}} + \mathcal{H}_{\text{SO}} \quad (3a)$$

$$\mathcal{H}_0 = \sum_{k,\alpha,s} \epsilon_{k,\alpha} c_{k,\alpha,s}^\dagger c_{k,\alpha,s} \quad (3b)$$

$$\mathcal{H}_Z = \Delta_Z \sum_{k,\alpha,s} s c_{k,\alpha,s}^\dagger c_{k,\alpha,s} \quad (3c)$$

$$\mathcal{H}_{\text{SO}} = \sum_{k,\alpha,\alpha',s,s'} L_{\alpha,\alpha',s,s'}(k) c_{k,\alpha,s}^\dagger c_{k,\alpha',s'} \quad (3d)$$

where $\alpha = 1$ (nearby), 2 (conduction) is the band index with $s = (\uparrow)$, (\downarrow) spin, $\epsilon_{k,\alpha} = \hbar^2 k^2 / 2m_\alpha^* - \delta_{\alpha,1}\Delta$ is the single-particle dispersion with $m_\alpha^* = (-1)^\alpha m^*$ effective mass and Δ band gap, $\Delta_Z = g\mu_B B_z$ is the Zeeman energy. $\mathcal{H}_{\text{scatt}}$ is responsible for the finite quasi-particle lifetime due to impurity and electron-phonon scattering and $L_{\alpha,\alpha',s,s'}(k)$ is the SOC.

The corresponding band structure is depicted in Fig. 2. The eigen-energies and eigenstates without SOC are

$$e_{k,\alpha,s} = \epsilon_{k,\alpha} + s\Delta_Z \quad (4a)$$

$$|1,\downarrow\rangle = [1,0,0,0]^\top \quad |1,\uparrow\rangle = [0,1,0,0]^\top \quad (4b)$$

$$|2,\downarrow\rangle = [0,0,1,0]^\top \quad |2,\uparrow\rangle = [0,0,0,1]^\top. \quad (4c)$$

The most general expression of the SOC for the above levels reads:

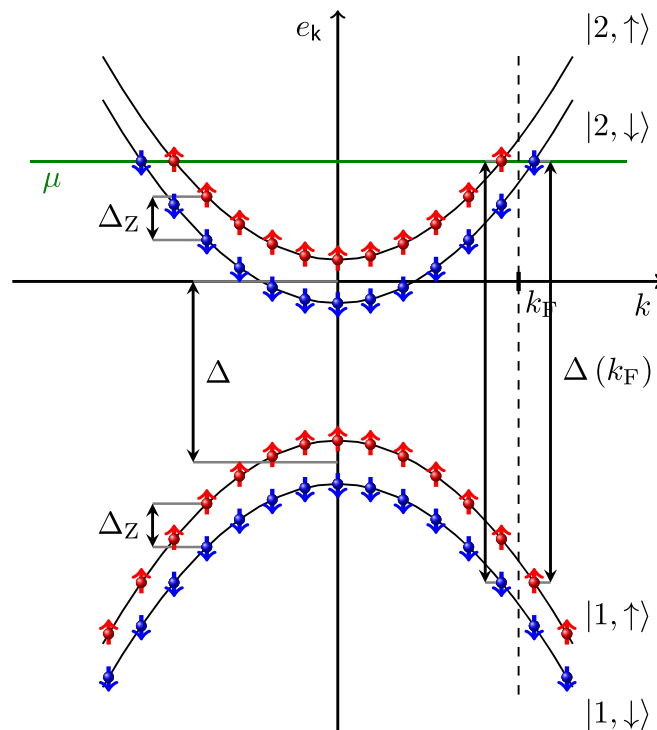


Figure 2 | The band structure of a 2DEG in a magnetic field. The effects of the weak SOC are not shown. Vertical arrows show the energy separations between the relevant bands.



$$L_{\alpha,\alpha',s,s'}(k) = \begin{pmatrix} \mathcal{L}\uparrow\uparrow & \mathcal{L}\uparrow\downarrow & L\uparrow\uparrow & L\uparrow\downarrow \\ \mathcal{L}\uparrow\downarrow & \mathcal{L}\downarrow\downarrow & L\uparrow\downarrow & L\downarrow\downarrow \\ L\uparrow\uparrow & L\uparrow\downarrow & \mathcal{L}\uparrow\uparrow & \mathcal{L}\uparrow\downarrow \\ L\uparrow\downarrow & L\downarrow\downarrow & \mathcal{L}\uparrow\downarrow & \mathcal{L}\downarrow\downarrow \end{pmatrix}, \quad (5)$$

where $\mathcal{L}_{s,s'}(k)$, $L_{ss'}(k)$ are the wavevector dependent intra- and inter-band terms, respectively, which are phenomenological, i.e. not related to a microscopic model. The terms mixing the same spin direction can be ignored as they commute with the S_z operator and do not cause spin-relaxation. The SOC terms contributing to spin-relaxation are

$$L_{\alpha,\alpha',s,s'}(k) = \begin{pmatrix} 0 & \mathcal{L} & 0 & L \\ \mathcal{L}^\dagger & 0 & L^\dagger & 0 \\ 0 & L & 0 & \mathcal{L} \\ L^\dagger & 0 & \mathcal{L}^\dagger & 0 \end{pmatrix}. \quad (6)$$

Table I. summarizes the role of the inversion symmetry on the SOC parameters. For a material with inversion symmetry, the Kramers theorem dictates (without magnetic field) that $\epsilon_\uparrow(k) = \epsilon_\downarrow(k)$ and thus $\mathcal{L} = 0$, which term would otherwise split the spin degeneracy in the same band. When the inversion symmetry is broken, \mathcal{L} is finite and the previous degeneracy is reduced to a weaker condition: $\epsilon_\uparrow(k) = \epsilon_\downarrow(-k)$ dictated by time reversal symmetry.

We consider the SOC as the smallest energy scale in our model ($\mathcal{L}(k_F)$, $L(k_F)$), while we allow for a competition of the other energy scales, namely Δ_Z , Γ and Δ , which can be of the same order of magnitude, as opposed to the conventional EY or DP case. We are mainly interested in the regime of a weak SOC, moderate magnetic fields, high occupation, and a large band gap. We treat the quasi-particle scattering rate to infinite order thus large values of Γ are possible.

The energy spectrum of the spins (or the ESR line-width) can be calculated from the Mori-Kawasaki formula^{27,28}, which relies on the assumption that the line-shape is Lorentzian. This was originally proposed for localized spins (e.g. Heisenberg-type models) but it can be extended to itinerant electrons. The standard (Faraday) ESR configuration measures the absorption of the electromagnetic wave polarized perpendicular to the static magnetic field. The ESR signal intensity is

$$I(\omega) = \frac{B_\perp^2 \omega}{2\mu_0} \chi''_\perp(q=0, \omega) V, \quad (7)$$

where B_\perp is the magnetic induction of the electromagnetic radiation, χ''_\perp is the imaginary part of the spin-susceptibility, μ_0 is the permeability of vacuum, and V is the sample volume. The spin-susceptibility is related to the retarded Green's function as

$$\chi''_\perp(\omega) = -\text{Im}G_{S^+S^-}^R(\omega), \quad (8)$$

with $S^\pm = S_x \pm iS_y$, from which the ESR spectrum can be obtained.

The equation of motion of the S^+ operator reads as

$$\frac{dS^+}{dt} = \frac{i}{\hbar} [\mathcal{H}, S^+] = \underbrace{\frac{i}{\hbar} [\mathcal{H}_Z, S^+]}_{-i\Delta_Z \frac{S^+}{\hbar}} + \underbrace{\frac{i}{\hbar} [\mathcal{H}_{\text{SO}}, S^+]}_{i\mathcal{A}}, \quad (9)$$

where $\mathcal{A} = \frac{1}{\hbar} [\mathcal{H}_{\text{SO}}, S^+]$ is the consequence of the SOC. The Green's function of S^+S^- is obtained from the Green's function of $\mathcal{A}^\dagger \mathcal{A}$ as

$$G_{S^+S^-}^R(\omega) = \frac{2\langle S_z \rangle}{\omega - \frac{\Delta_Z}{\hbar}} + \frac{-\langle [\mathcal{A}(0), S^-(0)] \rangle + G_{\mathcal{A}^\dagger \mathcal{A}}^R(\omega)}{\left(\omega - \frac{\Delta_Z}{\hbar}\right)^2}. \quad (10)$$

Table I | Effect of the presence or absence of the inversion symmetry on the intra- (\mathcal{L}) and inter-band (L) SOC and on the energy splitting of spin-states in the same band, $\epsilon_{k,\uparrow} - \epsilon_{k,\downarrow}$

	inversion symmetry	broken inv. symm.
L	0	finite
$\epsilon_{k,\uparrow} - \epsilon_{k,\downarrow}$	finite	finite
	0	finite

The second term is zero without SOC thus a completely sharp resonance occurs at the Zeeman energy. The line-shape is Lorentzian for a weak SOC:

$$G_{S^+S^-}^R(\omega) = \frac{2\hbar\langle S_z \rangle}{\hbar\omega - \Delta_Z - \Sigma(\omega)}, \quad (11)$$

where the self-energy is

$$\Sigma(\omega) = \frac{-\langle [\mathcal{A}(0), S^-(0)] \rangle + G_{\mathcal{A}^\dagger \mathcal{A}}^R(\omega)}{2\langle S_z \rangle}, \quad (12)$$

which is assumed to be a smooth function of ω near Δ_Z/\hbar .

The spin-relaxation rate is equal to the imaginary part of $\Sigma(\omega)$ as

$$\Gamma_s = \frac{\text{Im}G_{\mathcal{A}^\dagger \mathcal{A}}^R\left(\frac{\Delta_Z}{\hbar}\right)}{2\langle S_z \rangle}. \quad (13)$$

The $G_{\mathcal{A}^\dagger \mathcal{A}}^R(\omega)$ correlator is obtained from the Matsubara Green's function of $\mathcal{A}^\dagger \mathcal{A}$, given by

$$G_{\mathcal{A}^\dagger \mathcal{A}}(iv_m) = \int_0^{\beta\hbar} d\tau e^{iv_m\tau} \langle \mathcal{T}_\tau \mathcal{A}^\dagger(\tau) \mathcal{A}(0) \rangle. \quad (14)$$

where v_m is the bosonic Matsubara frequency. The effect of $\mathcal{H}_{\text{scatt}}$ is taken into account in the Green's function by a finite, constant momentum-scattering rate.

The most compact form of the spin-relaxation is obtained when the Fermi energy is not close to the bottom of the conduction band ($\mu \gtrsim \Delta$) and a calculation (detailed in the Methods section) using Eq. (13) leads to our main result:

$$\Gamma_s = \frac{4\Gamma|\mathcal{L}(k_F)|^2}{4\Gamma^2 + \Delta_Z^2} + \frac{4\Gamma|L(k_F)|^2}{4\Gamma^2 + \Delta^2(k_F)}, \quad (15)$$

Results in more general cases are discussed in the Supplementary Material.

Discussion

According to Eq. (15), the contributions from intra- ($\mathcal{L}(k_F)$) and inter-band ($L(k_F)$) processes are additive to lowest order in the SOC and have a surprisingly similar form. A competition is observed between lifetime induced broadening (due to Γ) and the energy separation between states ($\Delta(k_F)$ or Δ_Z). The situation, together with schematics of the corresponding band-structures, is shown in Fig. 3. When the broadening is much smaller than the energy separation, the relaxation is EY-like, $\Gamma_s \propto \Gamma$, even when the intra-band SOC dominates, i.e. for a material with inversion symmetry breaking. This situation was also studied in Ref. 29, 30 and it may be realized in III-V semiconductors in high magnetic fields. For metals with inversion symmetry, this is the canonical EY regime.

When the states are broadened beyond distinguishability (i.e. $\Gamma \gg \Delta(k_F)$ or Δ_Z), spin-relaxation is caused by two quasi-degenerate states and the relaxation is of DP-type, $\Gamma_s \propto 1/\Gamma$, even for a metal with inversion symmetry, $\mathcal{L} = 0$. For usual metals, the $\Gamma \gg \Delta(k_F)$,

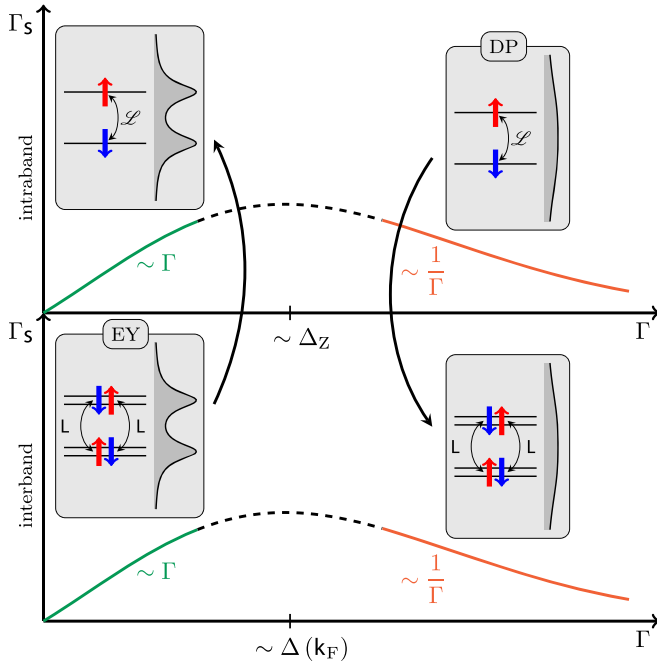


Figure 3 | Γ_s as a function of Γ according to Eq. (15) concerning separately the contributions due to intra- (upper) and inter-band (lower) SOC's. The insets are schematics of the band-structure including the broadening due to Γ (double arrows indicate the matrix elements). The arrows indicate the equivalence of the different spin-relaxation regimes. The conventional DP and EY scenarios are those in the top-right and bottom-left corners, respectively.

criterion implies a breakdown of the quasi-particle picture as therein $\Delta(k_F)$ is comparable to the bandwidth, thus this criterion means strong-localization. In contrast, metals with nearly degenerate bands remain metallic as e.g. MgB_2 (Ref. 25) and the alkali fullerenes (K_3C_{60} and Rb_3C_{60}) (Ref. 26), which are strongly correlated metals with large Γ . When the intra-band SOC dominates, i.e. for a strong inversion symmetry breaking, this is the canonical DP regime. These observations provide the ultimate link between these two spin-relaxation mechanisms, which are conventionally thought as being mutually exclusive.

Similar behavior can be observed in other models (details are given in the Supplementary material), $\Gamma_s \propto \Gamma$ and $\Gamma_s \propto 1/\Gamma$ remain valid in the two different limits but the intermediate behavior is not universal. A particularly compelling situation is the case of graphene where a four-fold degeneracy is present at the Dirac-point and both inter- and intra-band SOC are present thus changing the chemical potential would allow to map the crossovers predicted herein.

Methods

We consider Eq. (14) as a starting point. The Matsubara Green's function of $\mathcal{A}^\dagger \mathcal{A}$ can be written as

$$\mathcal{G}_{\mathcal{A}^\dagger \mathcal{A}}(iv_m) = \frac{1}{\beta \hbar} \sum_{i\omega_n, k, \alpha, s, s'} |\langle k, \alpha, s | \mathcal{A}^\dagger | k, \alpha', s' \rangle|^2 \mathcal{G}_{\alpha, s}(i\omega_n, k) \mathcal{G}_{\alpha', s'}(i\omega_n + iv_m, k), \quad (16)$$

where

$$\mathcal{G}_{\alpha, s}(i\omega_n, k) = \frac{1}{i\omega_n - \frac{1}{\hbar}(e_{k, \alpha, s} - \mu) + i \frac{\Gamma}{\hbar} \text{sgn}(\omega_n)} \quad (17)$$

is the Matsubara Green's function of fermionic field operators in band (α) and spin (s). The effect of $\mathcal{H}_{\text{scatt}}$ is taken into account by the finite momentum-scattering rate, Γ .

Using the relationship between the Green's function and spectral density, the Matsubara summation in Eq. (16) yields

$$\mathcal{G}_{\mathcal{A}^\dagger \mathcal{A}}(iv_m) = \frac{1}{4\pi^2} \sum_{k, \alpha, s, s'} |\langle k, \alpha, s | \mathcal{A}^\dagger | k, \alpha', s' \rangle|^2 \times \int_{-\infty}^{\infty} \int_{-\infty}^{\infty} d\omega d\omega' n_F(\omega'') \frac{n_F(\omega'') - n_F(\omega')}{i v_m - \omega'' + \omega'} \rho_{\alpha, s}(\omega', k) \rho_{\alpha', s'}(\omega'', k), \quad (18)$$

where

$$\rho_{\alpha, s}(\omega, k) = \frac{2\hbar\Gamma}{[\hbar\omega - (e_{k, \alpha, s} - \mu)]^2 + (\Gamma)^2} \quad (19)$$

is the spectral density. By taking the imaginary part after analytical continuation, the energy integrals can be calculated at zero temperature. Then, by replacing momentum summation with integration, we obtain

$$\text{Im} \mathcal{G}_{\mathcal{A}^\dagger \mathcal{A}}^R(\omega) = \frac{A}{8\pi^2} \int_0^\infty dk k \sum_{\alpha, \alpha', s, s'} |\langle k, \alpha, s | \mathcal{A}^\dagger | k, \alpha', s' \rangle|^2 \zeta_{\alpha, s, \alpha', s'}(k, \omega), \quad (20)$$

where

$$\zeta_{\alpha, s, \alpha', s'}(k, \omega) = \frac{4\hbar\Gamma}{4\Gamma^2 + (\tilde{e}_{k, \alpha, s} - \tilde{e}_{k, \alpha', s'} - \hbar\omega)^2} \times \left[\arctan\left(\frac{\tilde{e}_{k, \alpha, s}}{\Gamma}\right) - \arctan\left(\frac{\tilde{e}_{k, \alpha', s'}}{\Gamma}\right) - \arctan\left(\frac{\tilde{e}_{k, \alpha, s} - \hbar\omega}{\Gamma}\right) + \arctan\left(\frac{\tilde{e}_{k, \alpha', s'} + \hbar\omega}{\Gamma}\right) \right] + \frac{4\hbar\Gamma^2}{(\tilde{e}_{k, \alpha, s} - \tilde{e}_{k, \alpha', s'} - \hbar\omega) [4\Gamma^2 + (\tilde{e}_{k, \alpha, s} - \tilde{e}_{k, \alpha', s'} - \hbar\omega)^2]} \times \ln \frac{[\tilde{e}_{k, \alpha, s}^2 + \Gamma^2] [\tilde{e}_{k, \alpha', s'}^2 + \Gamma^2]}{[(\tilde{e}_{k, \alpha, s} - \hbar\omega)^2 + \Gamma^2] [(\tilde{e}_{k, \alpha', s'} + \hbar\omega)^2 + \Gamma^2]}$$

and $\tilde{e}_{k, \alpha, s} = e_{k, \alpha, s} - \mu$, A is the area of the 2DEG. The matrix elements of the \mathcal{A}^\dagger operator are

$$\langle k, \alpha, s | \mathcal{A}^\dagger | k, \alpha', s' \rangle = \begin{pmatrix} -\mathcal{L} & 0 & -L & 0 \\ 0 & \mathcal{L} & 0 & L \\ -L & 0 & -\mathcal{L} & 0 \\ 0 & L & 0 & \mathcal{L} \end{pmatrix}. \quad (21)$$

We determine the expectation value of the z-component of electron spin following similar steps as

$$\langle S_z \rangle = \sum_{i\omega_n, k, \alpha, s} \langle k, \alpha, s | S_z | k, \alpha, s \rangle \mathcal{G}_{\alpha, s}(i\omega_n, k) = \sum_{k, \alpha, s} \langle k, \alpha, s | S_z | k, \alpha, s \rangle \int_{-\infty}^{\infty} \frac{d\omega'}{2\pi} n_F(\omega') \rho_{\alpha, s}(\omega', k) = \sum_{k, \alpha, s} \langle k, \alpha, s | S_z | k, \alpha, s \rangle \zeta_{\alpha, s}(k), \quad (22)$$

where

$$\zeta_{\alpha, s}(k) = \frac{1}{2} - \frac{1}{\pi} \arctan\left(\frac{\tilde{e}_{k, \alpha, s}}{\Gamma}\right). \quad (23)$$

The matrix elements of the S_z operator are

$$\langle k, \alpha, s | S_z | k, \alpha, s \rangle = \frac{\hbar}{2} \begin{pmatrix} -1 & 0 & 0 & 0 \\ 0 & 1 & 0 & 0 \\ 0 & 0 & -1 & 0 \\ 0 & 0 & 0 & 1 \end{pmatrix}. \quad (24)$$

The spin-relaxation rate can be obtained as

$$\Gamma_s = \frac{\text{Im} \mathcal{G}_{\mathcal{A}^\dagger \mathcal{A}}^R\left(\frac{\Delta Z}{\hbar}\right)}{2\langle S_z \rangle} = \Gamma_s^{\text{intra}} + \Gamma_s^{\text{inter}}. \quad (25)$$

We note this is the sum of intra- and inter-band terms which are described separately.

1. Wolf, S. A. *et al.* Spintronics: A spin-based electronics vision for the future. *Science* **294**, 1488–1495 (2001).
2. Žutić, I., Fabian, J. & Sarma, S. D. Spintronics: Fundamentals and applications. *Rev. Mod. Phys.* **76**, 323–410 (2004).
3. Wu, M. W., Jiang, J. H. & Weng, M. Q. Spin dynamics in semiconductors. *Phys. Rep.* **493**, 61–236 (2010).
4. Feher, G. & Kip, A. F. Electron Spin Resonance Absorption in Metals. I. Experimental. *Physical Review* **98**, 337–348 (1955).
5. Johnson, M. & Silsbee, R. H. Coupling of electronic charge and spin at a ferromagnetic-paramagnetic metal interface. *Phys. Rev. B* **37**, 5312–5325 (1988).



6. Jedema, F., Heersche, H., Filip, A., Baselmans, J. & van Wees, B. Electrical detection of spin precession in a metallic mesoscopic spin valve. *Nature* **416**, 713–716 (2002).
7. Huertas-Hernando, D., Guinea, F. & Brataas, A. Spin-orbit coupling in curved graphene, fullerenes, nanotubes, and nanotube caps. *Phys. Rev. B* **74**, 155426 (2006).
8. Ertler, C., Kunschuh, S., Gmitra, M. & Fabian, J. Electron spin relaxation in graphene: the role of the substrate. *Phys. Rev. B* **80**, 041405 (2009).
9. Gmitra, M., Kunschuh, S., Ertler, C., Ambrosch-Draxl, C. & Fabian, J. Band-structure topologies of graphene: spin-orbit coupling effects from first principles. *Phys. Rev. B* **80**, 235431 (2009).
10. Castro Neto, A. H. & Guinea, F. Impurity-induced spin-orbit coupling in graphene. *Phys. Rev. Lett.* **103**, 026804 (2009).
11. Dóra, B., Murányi, F. & Simon, F. Electron spin dynamics and electron spin resonance in graphene. *Eur. Phys. Lett.* **92**, 17002 (2010).
12. Zhang, P. & Wu, M. W. Electron spin relaxation in graphene with random Rashba field: comparison of the D'yakonov-Perel' and Elliott-Yafet-like mechanisms. *New J. Phys.* **14**, 033015 (2012).
13. Ochoa, H., Castro Neto, A. H. & Guinea, F. Elliot-yafet mechanism in graphene. *Phys. Rev. Lett.* **108**, 206808 (2012).
14. Tombros, N., Józsa, C., Popinciuc, M., Jonkman, H. T. & van Wees, B. J. Electronic spin transport and spin precession in single graphene layers at room temperature. *Nature* **448**, 571–574 (2007).
15. Han, W. *et al.* Tunneling Spin Injection into Single Layer Graphene. *Phys. Rev. Lett.* **105**, 167202 (2010).
16. Han, W. & Kawakami, R. K. Spin relaxation in single-layer and bilayer graphene. *Phys. Rev. Lett.* **107**, 047207 (2011).
17. Yang, T.-Y. *et al.* Observation of long spin-relaxation times in bilayer graphene at room temperature. *Phys. Rev. Lett.* **107**, 047206 (2011).
18. Pikus, G. E. & Titkov, A. N. In Meier, F. & Zakharchenya, B. (eds.) *Optical Orientation* (North-Holland, Amsterdam, 1984).
19. Averkiev, N., Golub, L. & Willander, M. Spin relaxation anisotropy in two-dimensional semiconductor systems. *J. Phys. Cond. Mat.* **14**, R271–R283 (2002).
20. Glazov, M. M., Sherman, E. Y. & Dugaev, V. K. Two-dimensional electron gas with spin-orbit coupling disorder. *Phys. E* **42**, 2157–2177 (2010).
21. Elliott, R. J. Theory of the Effect of Spin-Orbit Coupling on Magnetic Resonance in Some Semiconductors. *Phys. Rev.* **96**, 266–279 (1954).
22. Yafet, Y. Conduction electron spin relaxation in the superconducting state. *Physics Letters A* **98**, 287–290 (1983).
23. Dyakonov, M. & Perel, V. Spin relaxation of conduction electrons in noncentrosymmetric semiconductors. *Soviet Physics Solid State, USSR* **13**, 3023–3026 (1972).
24. Tombros, N. *et al.* Anisotropic spin relaxation in graphene. *Phys. Rev. Lett.* **101**, 046601-1–4 (2008).
25. Simon, F. *et al.* Generalized Elliott-Yafet Theory of Electron Spin Relaxation in Metals: Origin of the Anomalous Electron Spin Lifetime in MgB₂. *Phys. Rev. Lett.* **101**, 177003-1–4 (2008).
26. Dóra, B. & Simon, F. Electron-spin dynamics in strongly correlated metals. *Phys. Rev. Lett.* **102**, 137001 (2009).
27. Mori, H. & Kawasaki, K. Antiferromagnetic resonance absorption. *Progress of Theoretical Physics* **28**, 971–987 (1962).
28. Oshikawa, M. & Affleck, I. Electron spin resonance in $s = \frac{1}{2}$ antiferromagnetic chains. *Phys. Rev. B* **65**, 134410 (2002).
29. Ivchenko, E. L. Spin relaxation of free carriers in a noncentrosymmetric semiconductor in a longitudinal magnetic field. *Sov. Phys. Solid State* **15**, 1048 (1973).
30. Burkov, A. A. & Balents, L. Spin relaxation in a two-dimensional electron gas in a perpendicular magnetic field. *Phys. Rev. B* **69**, 245312 (2004).

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Author contributions

P.B. carried out all calculations under the guidance of B.D. A.K. contributed to the discussion and F.S. initiated the development of the unified theory. All authors contributed to the writing of the manuscript.

Additional information

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