Spin Hall Conductance In a Two-Dimensional Tight-Binding Model In The Presence Of Rashba Spin-Orbit Interaction And Random Impurities

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Abstract

The spin-Hall conductance in a two-dimensional tight-binding model is studied in the presence of Rashba spin-orbit interaction and random impurities. The conventional definition of spin current falls through for a system with Rashba spin-orbit interaction because of the non-conservation of the spin magnetic moment in the presence of Rashba coupling. Using a modified definition of spin-current operator, the relaxation time for electron-impurity scattering is calculated with the help of Matsubara Green function and the spin-Hall conductivity is determined by employing the Kubo-Greenwood formalism. Our results show the explicit behavior of the spin-Hall conductance as a function of chemical potential, Rashba spin-orbit coupling constant and the impurity strength

Keywords: *Rashba Spin-Orbit Interaction, Random Impurities, Kubo Greenwood formalism, Spin Hall Conductivity, Relaxation Time*

1. Introduction

Recent years have witnessed a flurry of investigations on the subject of spin transport in low-dimensional systems. In this connection, several new ideas have been mooted for the realization of new devices that would mainly rely on the spin degrees of freedom of the electrons [1,3]. Of late, both intrinsic and extrinsic spin-Hall effect [5-8] have attracted considerable attention. However majority of the works reported so far have used the conventional definition of spin-current $(\frac{1}{2})$ $\frac{1}{2} [v s^z + s^z v]$, which unfortunately falls through in the presence of Rashba spin-orbit interaction (RSOI) because the spin magnetic moment of the system is not conserved in the presence of the Rashba coupling. Furthermore, the conventional definition of spin current gives a finite result in insulators even with localized states. To get rid of the afore-mentioned difficulties, we have constructed following the definition suggested by Shi et al. [1] a modified spin-current operator for our two-dimensional tight-binding model with spin-orbit coupling in the presence of impurity.

 We consider a two-dimensional electron system with RSOI in the presence of impurity within the framework of the tight-binding model. The Hamiltonian for the present system can be written as

$$
H = H_0 + H_{int} \quad , \tag{1}
$$

with

$$
H_0 = \varepsilon_0 \sum_i c_i^{\dagger} c_i + t \sum_{\langle ij \rangle} (c_i^{\dagger} c_j + h.c) - i \alpha_R \sum_{\langle i,j \rangle} (c_{ix,iy}^{\dagger} \sigma^y c_{ix+1,iy} + h.c) + i \alpha_R \sum_{\langle i,j \rangle} (c_{ix,iy}^{\dagger} \sigma^x c_{ix,iy+1} + h.c), \tag{2}
$$

$$
H_{int} = \sum \epsilon_i \, c_i^{\dagger} c_i = \sum_{i,l} v \delta(r_i - r_l) c_i^{\dagger} c_i \tag{3}
$$

where ε_0 is the site energy, t is the hopping integral, α_R is the Rashba spin-orbit coupling constant, v is the impurity potential strength, $(\sigma_x, \sigma_y, \sigma_z)$ are the Pauli matrices and

$$
c_i = \begin{pmatrix} c_{i\uparrow} \\ c_{i\downarrow} \end{pmatrix} , \quad c_i^{\dagger} = \begin{pmatrix} c_{i\uparrow}^{\dagger} & c_{i\downarrow}^{\dagger} \end{pmatrix} . \tag{4}
$$

 $c_{i\uparrow j\downarrow}^{\dagger}(c_{i\uparrow j\downarrow})$ is the creation (annihilation) operator for a spin-up/spin-down electron at the *i*-th site. In the presence of Rashba spin-orbit interaction, the spin degeneracy is lifted leading to two non-degenerate bands. When the impurity is present, the charge carriers will acquire a finite relaxation time τ through elastic scattering. This relaxation time can be calculated from the imaginary part of the self-energy.

2. Theoretical Formalism

2.1. Spin and Charge currents

We define the spin polarization operator as

$$
\vec{P}^{s_z} = \sum_{x_i, y_i} \vec{R}_{x_i, y_i} c^{\dagger}_{x_i, y_i} \sigma^z , \qquad (5)
$$

so that the spin current can be written as

$$
J^{s_z} = \frac{\partial \vec{P}^{s_z}}{\partial t} = i \left[H \sum_{x_i, y_i} \vec{R}_{x_i, y_i} c_{x_i, y_i}^\dagger \sigma^z c_{x_i, y_i} \right] \tag{6}
$$

Similarly, the charge polarization and the charge current can be written as

$$
\vec{P}^c = \sum_{x_i, y_i} \vec{R}_{x_i, y_i} c_{x_i, y_i}^\dagger I c_{x_i, y_i} \quad , \tag{7}
$$

$$
J^{c} = \frac{\partial \vec{P}^{c}}{\partial t} = i \left[H \sum_{x_{i}, y_{i}} \vec{R}_{x_{i}, y_{i}} c_{x_{i}, y_{i}}^{\dagger} l c_{x_{i}, y_{i}} \right] . \tag{8}
$$

2.2 Spin Hall Conductivity

According to the Kubo formalism, the spin-Hall conductivity is given by

$$
\sigma_{xy}^{S_Z} = -\frac{e\hbar}{\pi} \sum_{M \neq N} \frac{Im[]}{(E_M - E_N)^2 + \left(\frac{\hbar}{\tau(\mu)}\right)^2} (f_F(E_M) - f_F(E_n)),
$$
\n(9)

where $|M > |N >$ are the eigen states of H_0 belonging to the eigenvalues E_M and E_N respectively and $\tau(\mu)$ is the configuration-averaged relaxation time of the electron. The spin-Hall conductivity is calculated by substituting Eqs. (6) and (8) in Eq. (9)

2. Numerical results

We first calculate the spin and charge currents and the configuration-average relaxation times for the upspin and down-spin electrons using the Mastubara Green functions. The relaxation times for the up and down-spin electrons turn out to be equal. The spin-Hall conductivity is finally obtained by using the Kubo-Greenwood formula. Fig (1) shows the variation of spin-Hall conductivity (SHE) as a function of α_R for different values of v. Below a certain critical value of α_R (say α_{RC}) (which depends on v), SHC remanis vanishingly snall and as α_R exceeds α_{RC} , SHE increases with α_R motonically and quite rapidly. Fig. 2 presents the behaviour of SHC as a function of v for different values of μ . As expected, SHC decreases with increasing ν and becomes zero above a certain value of ν . The figure also shows that at small ν , SHC increases with the chemical potential.

Fig.1 Spin Hall Conductance vs RSOI strength Fig.2 Spin Hall Conductance vs Impurity

4. Conclusion

Using a modified definition of spin-current operator, SHC is calculated using the Matsubara Green function technique and the Kubo-Greenwood formula. It is shown that below a critical value of α_R , SHC is zero and above this critical α_R , SHC increases monotonically and rapidly. As a function of the impurity strength v , SHC, as expected, decreases quite rapidly with increasing v . At small values of ν , SHC turns out to be larger for smaller values of the chemical potential.

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