

Scanning tunneling microscope operating as a spin diode

P. H. Penteado,^{1,2} F. M. Souza,² A. C. Seridonio,³ E. Vernek,² and J. C. Egues¹

¹*Instituto de Física de São Carlos, Universidade de São Paulo, 13560-970, São Carlos, SP, Brazil*

²*Instituto de Física, Universidade Federal de Uberlândia, 38400-902, Uberlândia, MG, Brazil*

³*Departamento de Física e Química, Universidade Estadual Paulista Júlio de Mesquita Filho, 15385-000, Ilha Solteira, SP, Brazil.*

(Received 21 June 2011; published 19 September 2011)

We theoretically investigate spin-polarized transport in a system composed of a ferromagnetic scanning-tunneling-microscope (STM) tip coupled to an adsorbed atom (adatom) on a host surface. Electrons can tunnel directly from the tip to the surface or via the adatom. Since the tip is ferromagnetic and the host surface (metal or semiconductor) is nonmagnetic we obtain a spin-diode effect when the adatom is in the regime of single occupancy. This effect leads to an unpolarized current for direct bias ($V > 0$) and polarized current for reverse ($V < 0$) bias voltages, if the tip is nearby the adatom. Within the nonequilibrium Keldysh technique we analyze the interplay between the lateral displacement of the tip and the intra adatom Coulomb interaction on the spin-diode effect. As the tip moves away from the adatom the spin-diode effect vanishes and the currents become polarized for both $V > 0$ and $V < 0$. We also find an imbalance between the up and down spin populations in the adatom, which can be tuned by the tip position and the bias. Finally, due to the presence of the adsorbate on the surface, we observe spin-resolved Friedel oscillations in the current, which reflects the oscillations in the calculated local density of states (LDOS) of the subsystem surface + adatom.

DOI: [10.1103/PhysRevB.84.125439](https://doi.org/10.1103/PhysRevB.84.125439)

PACS number(s): 68.37.Ef, 75.76.+j, 73.23.Hk

I. INTRODUCTION

The scanning-tunneling microscope (STM) has allowed huge advances in condensed matter physics. On one hand it serves as a powerful tool to manipulate matter on a single atomic scale;^{1,2} on the other it is used as a probe of the topology of metallic and semiconductor surfaces.³ An impressive early example of such control is the quantum corral, assembled by moving atom-by-atom on a metallic surface.^{4,5}

In the fascinating field of spintronics,⁶ STM was recently used to manipulate individual Co atoms adsorbed on a template of Mn.⁷ It was possible, for instance, to determine the spin direction of the individual Co atoms. STM was also applied to study the interactions between isolated Mn acceptors and the influence of the surface on the impurity properties in diluted magnetic semiconductors, e.g., Mn-doped GaAs.^{8,9} More interesting, spin-polarized STM, sensitive to surface magnetization,¹⁰ has been used to map the morphology and the density of states of single magnetic structures¹¹ and magnetic quantum dots.¹² STM has also been employed in the investigation and identification of promising molecular switches, which could be used in future nanoscale circuits.^{13–15} In the context of quantum information,¹⁶ STM was used to measure electron spin-relaxation times of individual atoms adsorbed on a surface with nanosecond time resolution.^{17,18} More recently, a new type of scanning probe microscope was demonstrated using ultracold atoms.¹⁹ Interestingly, the conventional solid tip is replaced by a gas of ultracold rubidium atoms, which increases the spatial resolution of the microscope. All these applications highlight the importance of STM to the development of nanoengineered systems for spintronics and spin-based quantum information processing.

As for strong correlated systems, STM has revealed new facets of the Kondo physics, such as the Fano-Kondo resonance on the differential conductance when a single magnetic adatom (e.g., Co) is placed on a metallic surface.^{5,20,21} If the tip is ferromagnetic the Fano-Kondo line shape becomes spin-

dependent²² and the setup can be used as a powerful spin filter.²³

Here, we study spin-dependent transport in a system composed of a ferromagnetic (FM) STM tip coupled to both an adsorbed atom and a host nonmagnetic (NM) surface. This geometry resembles a junction NM-QD-FM, where “QD” stands for quantum dot. In the present system, however, we have one additional ingredient: the tip can move laterally away from the adatom. It is well known that the NM-QD-FM system gives rise to unpolarized current for direct bias ($V > 0$) and polarized current for reverse bias ($V < 0$) when the dot is singly occupied (Coulomb blockade forbids double occupancy).²⁴ This rectification of the current polarization is the so-called spin-diode effect and has been observed in carbon nanotubes.²⁵ In this work we investigate the interplay between

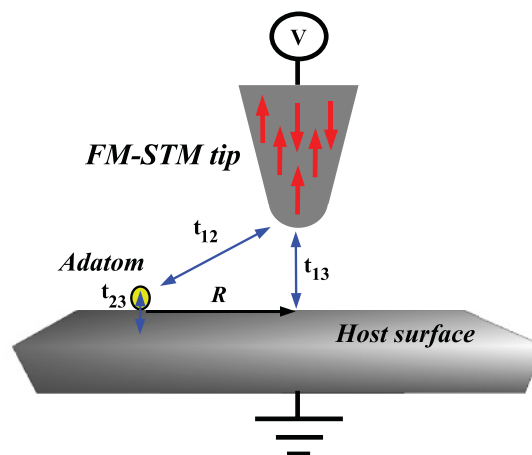


FIG. 1. (Color online) Ferromagnetic scanning-tunneling-microscope (FM-STM) tip coupled to a host surface with an adatom. The matrix elements t_{12} , t_{13} , and t_{23} represent tip-adatom, tip-surface, and adatom-surface couplings, respectively. The tip-adatom lateral distance is denoted by R .

the Coulomb blockade and the lateral displacement of the tip on the spin-diode effect.

Our calculation is based on the Keldysh nonequilibrium technique. By performing a self-consistent calculation we determine the adatom occupation and its magnetization as a function of the tip lateral position R , see Fig. 1. We find that the adatom becomes magnetized when the tip approaches it; this magnetization switches sign as the bias is reversed. Moreover, we calculate the spin-resolved current in both single- and double-occupancy regimes of the adatom and find that it can depend strongly on the tip position. In particular, the spin-diode effect emerges when the tip is closer to the adatom and the charging energy is large enough to allow for only a single electron in the adatom. As the tip moves away from the adatom the currents become equally polarized for both biases thus resulting in a suppression of the spin-diode behavior. More interesting, we observe spin-resolved Friedel oscillations^{26–30} in the current as the lateral distance tip-adatom R increases. These are due to the presence of the adatom on the surface, and reflect the oscillations in the local density of states (LDOS) of the subsystem surface + adatom.

The paper is organized as follows. In Sec. II we present a detailed description of the formulation used to compute the spin-resolved currents and the spin populations. We divide this section into three subsections: in A, we determine an expression for the spin-resolved currents and the local density of states of the subsystem adatom-surface when the tip is far away from the adatom. In B, the calculation for the current is performed in the presence of the tip, and in C we summarize the numerical technique and the parameters used in the simulations. We present our results and discussions in Sec. III. Section IV summarizes the main ideas of our work.

II. FORMULATION

The system we study is composed of a FM-STM tip over an adsorbate on a host surface, see Fig. 1. Electrons can tunnel from the tip to the substrate or vice-versa via two possible ways: (i) direct tunneling tip-to-surface or (ii) tunneling via the localized state of the adatom. The system Hamiltonian is

$$H = H_1 + H_2 + H_3 + H_{12} + H_{13} + H_{23}, \quad (1)$$

where H_i corresponds to the tip ($i = 1$), the adatom ($i = 2$), or the surface ($i = 3$), and reads

$$H_i = \sum_{\mathbf{k}_i\sigma} \epsilon_{\mathbf{k}_i\sigma} c_{\mathbf{k}_i\sigma}^\dagger c_{\mathbf{k}_i\sigma} + \delta_{i2} U \hat{n}_\uparrow \hat{n}_\downarrow, \quad (2)$$

where for $i = 1(3)$, \mathbf{k}_i is the wave number for electrons in the tip (host) and the label σ stands for the electron spin components \uparrow and \downarrow . Here, $\epsilon_{\mathbf{k}_i\sigma}$ is the energy of the state $\mathbf{k}_i\sigma$ and $c_{\mathbf{k}_i\sigma}$ ($c_{\mathbf{k}_i\sigma}^\dagger$) annihilates (creates) an electron in the quantum state $\mathbf{k}_i\sigma$. We consider a Stoner-like ferromagnetic dispersion $\epsilon_{\mathbf{k}_1\sigma} = \hbar^2 k_1^2 / 2m + \sigma \Delta$ for the tip, with m being the free-electron mass and Δ the usual Stoner parameter,^{31,32} and a free-electron dispersion $\epsilon_{\mathbf{k}_3} = \hbar^2 k_3^2 / 2m$ for the surface. For the adatom, $i = 2$, we consider only a single spin-degenerate energy level, $\epsilon_{\mathbf{k}_2\sigma} = \epsilon_\sigma$. In this case, the index \mathbf{k}_2 simply denotes the adatom level. The second term in H_2 accounts for the Coulomb interaction U in the adatom.

The coupling terms in Eq. (1) can be written as

$$H_{ij} = \sum_{\mathbf{k}_i, \mathbf{k}_j, \sigma} (t_{ij} c_{\mathbf{k}_i, \sigma}^\dagger c_{\mathbf{k}_j, \sigma} + t_{ij}^* c_{\mathbf{k}_j, \sigma}^\dagger c_{\mathbf{k}_i, \sigma}), \quad (3)$$

where t_{ij} is the coupling parameter between subsystems i and j ; t_{12} , t_{13} , and t_{23} account for the tip-adatom, tip-surface, and adatom-surface tunnelings, respectively. When a bias voltage is applied, these transfer terms drive the system out of equilibrium.

Next, we consider a real-space formulation for the spin-resolved current. This is particularly convenient since we are interested in looking at Friedel oscillations on the surface. As we shall see later on, this formulation is equivalent to a formulation in the \mathbf{k} space.

A. Nonresonant transport

For simplicity, let us first consider the transport regime in which the direct coupling between the tip and the adatom is negligible (nonresonant transport), which is valid for large enough tip-adatom lateral distances. The Hamiltonian of the system in this case reduces to

$$H = H_1 + H_2 + H_3 + H_{13} + H_{23}. \quad (4)$$

The electrical current for spin σ between the tip and the surface can be calculated from the definition³³

$$I_1^\sigma = -e \langle \dot{N}_1^\sigma \rangle = -ie \langle [H, N_1^\sigma] \rangle, \quad (5)$$

with e the electron charge ($e > 0$) and N_1^σ the total number operator given by

$$N_1^\sigma = \int d\mathbf{r}_1 \Psi_1^{\sigma\dagger}(\mathbf{r}_1, t) \Psi_1^\sigma(\mathbf{r}_1, t), \quad (6)$$

where $\Psi_1^\sigma(\mathbf{r}_1, t)$ and $\Psi_1^{\sigma\dagger}(\mathbf{r}_1, t)$ are quantum field operators for the electrons in the tip. In Eq. (5) and throughout the paper we assume $\hbar = 1$.

The quantity $\langle O(t) \rangle$ defines the nonequilibrium average value of a physical observable denoted by the operator $O(t)$, and it is given by^{33,34}

$$\langle O(t) \rangle = \text{Tr}[\rho O(t)], \quad (7)$$

where ρ is the thermal equilibrium density matrix, $\rho = (\text{Tr} e^{-\beta H_0})^{-1} e^{-\beta H_0}$, with H_0 being the Hamiltonian containing only the H_i terms in Eq. (4), and $O(t)$ is in the Heisenberg picture, i.e., its time dependence is governed by the full Hamiltonian of Eq. (4).

The only nonvanishing term in the commutator of Eq. (5) is $[H_{13}, N_1^\sigma]$. For electrons with spin σ , the tip-surface coupling can be written as³⁵

$$H_{13} = \sum_\sigma \iint d\mathbf{r}_1 d\mathbf{r}_3 [T(\mathbf{r}_1, \mathbf{r}_3) \Psi_1^{\sigma\dagger}(\mathbf{r}_1, t) \Psi_3^\sigma(\mathbf{r}_3, t) + \text{H.c.}], \quad (8)$$

where $T(\mathbf{r}_1, \mathbf{r}_3)$ is a matrix element that accounts for the coupling between the tip and the surface, and $\Psi_3^\sigma(\mathbf{r}_3, t)$ is the quantum field operator for electrons in the surface. Calculating

$[H_{13}, N_1^\sigma]$ and using the result in Eq. (5) we find for the spin-resolved current

$$I_1^\sigma = ie \int \int d\mathbf{r}_1 d\mathbf{r}_3 [T(\mathbf{r}_1, \mathbf{r}_3) \langle \Psi_1^{\sigma\dagger}(\mathbf{r}_1, t) \Psi_3^\sigma(\mathbf{r}_3, t) \rangle - T^*(\mathbf{r}_1, \mathbf{r}_3) \langle \Psi_3^{\sigma\dagger}(\mathbf{r}_3, t) \Psi_1^\sigma(\mathbf{r}_1, t) \rangle]. \quad (9)$$

Defining the lesser Green function

$$G_\sigma^<(\mathbf{r}_3, t_3; \mathbf{r}_1, t_1) = i \langle \Psi_1^{\sigma\dagger}(\mathbf{r}_1, t_1) \Psi_3^\sigma(\mathbf{r}_3, t_3) \rangle, \quad (10)$$

we can rewrite the current as

$$I_1^\sigma = 2e \text{Re} \left[\int \int d\mathbf{r}_1 d\mathbf{r}_3 T(\mathbf{r}_1, \mathbf{r}_3) G_\sigma^<(\mathbf{r}_3, t; \mathbf{r}_1, t) \right]. \quad (11)$$

We now aim at determining $G_\sigma^<(\mathbf{r}_3, t; \mathbf{r}_1, t)$ in Eq. (11). To this end, we use the nonequilibrium Keldysh formalism. Similarly to the equilibrium case, here we introduce an ordered Green function

$$G_\sigma(\mathbf{r}_3, \tau_3; \mathbf{r}_1, \tau_1) = -i \langle T_C \Psi_3^\sigma(\mathbf{r}_3, \tau_3) \Psi_1^{\sigma\dagger}(\mathbf{r}_1, \tau_1) \rangle, \quad (12)$$

with the τ 's defined, however, on a contour C in the complex plane. The operator T_C , called contour-ordering operator, orders the operators according to the position of their time arguments on the contour. From the contour-ordered Green function, we can obtain the lesser $G_\sigma^<$, greater $G_\sigma^>$, retarded G_σ^r , and advanced G_σ^a Green functions, which are directly linked to the observables.

To obtain $G_\sigma^<$, $G_\sigma^>$, G_σ^r , and G_σ^a , we first determine the equation of motion for the Green function in Eq. (12),

$$\begin{aligned} & \left(i \frac{\partial}{\partial \tau_1} - \frac{\nabla_{\mathbf{r}_1}^2}{2m} \right) G_\sigma(\mathbf{r}_3, \tau_3; \mathbf{r}_1, \tau_1) \\ &= - \int d\mathbf{r}'_3 T(\mathbf{r}'_3, \mathbf{r}_1) G_\sigma(\mathbf{r}_3, \tau_3; \mathbf{r}'_3, \tau_1), \end{aligned} \quad (13)$$

or in the integral form

$$\begin{aligned} G_\sigma(\mathbf{r}_3, \tau_3; \mathbf{r}_1, \tau_1) &= \int \int d\mathbf{r}'_1 d\mathbf{r}'_3 \int_C d\tilde{\tau} G_\sigma(\mathbf{r}_3, \tau_3; \mathbf{r}'_3, \tilde{\tau}) \\ &\quad \times T(\mathbf{r}'_3, \mathbf{r}'_1) g_\sigma(\mathbf{r}'_1, \tilde{\tau}; \mathbf{r}_1, \tau_1), \end{aligned} \quad (14)$$

where g_σ is the free-electron Green function of the tip and the time integral is over the contour C ; then we perform an appropriate analytical continuation. This procedure consists essentially in replacing the contour integral over τ in Eq. (14) by a real-time integral over t . Here we follow the Langreth procedure.³⁶ For the lesser Green function $G_\sigma^<$, we have

$$\begin{aligned} G_\sigma^<(\mathbf{r}_3, t_3; \mathbf{r}_1, t_1) &= \int \int d\mathbf{r}'_1 d\mathbf{r}'_3 \int d\tilde{t} \\ &\quad \times [G_\sigma^r(\mathbf{r}_3, t_3; \mathbf{r}'_3, \tilde{t}) T(\mathbf{r}'_3, \mathbf{r}'_1) g_\sigma^<(\mathbf{r}'_1, \tilde{t}; \mathbf{r}_1, t_1) \\ &\quad + G_\sigma^<(\mathbf{r}_3, t_3; \mathbf{r}'_3, \tilde{t}) T(\mathbf{r}'_3, \mathbf{r}'_1) g_\sigma^a(\mathbf{r}'_1, \tilde{t}; \mathbf{r}_1, t_1)]. \end{aligned} \quad (15)$$

In the above equation, g_σ^a and $g_\sigma^<$ correspond to the analytically continued free-electron advanced and lesser Green functions of the tip, respectively. Throughout the paper we use lower case to denote the free-electron Green functions of the tip, the adatom, and the surface. We note that $G_\sigma^<(\mathbf{r}_3, t_3; \mathbf{r}_1, t_1)$ is coupled to $G_\sigma^r(\mathbf{r}_3, t_3; \mathbf{r}'_3, \tilde{t})$ and also to $G_\sigma^<(\mathbf{r}_3, t_3; \mathbf{r}'_3, \tilde{t})$. To completely determine $G_\sigma^<(\mathbf{r}_3, t_3; \mathbf{r}_1, t_1)$, we then need to

perform an iterative process and obtain a system of equations for the Green functions G_σ^r and $G_\sigma^<$.

Substituting Eq. (15) into Eq. (11) we obtain

$$\begin{aligned} I_1^\sigma &= 2e \text{Re} \left\{ \int \int \int \int d\mathbf{r}_1 d\mathbf{r}_3 d\mathbf{r}'_1 d\mathbf{r}'_3 d\tilde{t} T(\mathbf{r}_1, \mathbf{r}_3) T(\mathbf{r}'_3, \mathbf{r}'_1) \right. \\ &\quad \times [G_\sigma^r(\mathbf{r}_3, t; \mathbf{r}'_3, \tilde{t}) g_\sigma^<(\mathbf{r}'_1, \tilde{t}; \mathbf{r}_1, t) \\ &\quad \left. + G_\sigma^<(\mathbf{r}_3, t; \mathbf{r}'_3, \tilde{t}) g_\sigma^a(\mathbf{r}'_1, \tilde{t}; \mathbf{r}_1, t) \right\}. \end{aligned} \quad (16)$$

Performing a Fourier transform in the time coordinate we find

$$\begin{aligned} I_1^\sigma &= 2e \int \frac{d\omega}{2\pi} \text{Re} \left\{ \int \int \int \int d\mathbf{r}_1 d\mathbf{r}_3 d\mathbf{r}'_1 d\mathbf{r}'_3 T(\mathbf{r}_1, \mathbf{r}_3) T(\mathbf{r}'_3, \mathbf{r}'_1) \right. \\ &\quad \times [G_\sigma^r(\mathbf{r}_3, \mathbf{r}'_3, \omega) g_\sigma^<(\mathbf{r}'_1, \mathbf{r}_1, \omega) \\ &\quad \left. + G_\sigma^<(\mathbf{r}_3, \mathbf{r}'_3, \omega) g_\sigma^a(\mathbf{r}'_1, \mathbf{r}_1, \omega) \right\}. \end{aligned} \quad (17)$$

Additionally, Fourier transforming $g_\sigma^<$ and g_σ^a results in

$$\begin{aligned} I_1^\sigma &= 2e \int \frac{d\omega}{2\pi} \text{Re} \left\{ \sum_{\mathbf{k}_1} \int \int \int \int d\mathbf{r}_1 d\mathbf{r}_3 d\mathbf{r}'_1 d\mathbf{r}'_3 \right. \\ &\quad \times e^{-i\mathbf{k}_1 \cdot \mathbf{r}_1} T(\mathbf{r}_1, \mathbf{r}_3) e^{i\mathbf{k}_1 \cdot \mathbf{r}'_1} T(\mathbf{r}'_3, \mathbf{r}'_1) \\ &\quad \left. \times [G_\sigma^r(\mathbf{r}_3, \mathbf{r}'_3, \omega) g_{\mathbf{k}_1\sigma}^<(\omega) + G_\sigma^<(\mathbf{r}_3, \mathbf{r}'_3, \omega) g_{\mathbf{k}_1\sigma}^a(\omega)] \right\}. \end{aligned} \quad (18)$$

Now we assume that the coupling between the tip and the surface is local^{35,37} (point source), i.e., $T(\mathbf{r}_1, \mathbf{r}_3) = T^0(\mathbf{r}_1) \delta(\mathbf{r}_3 - \mathbf{R})$, $T^0(\mathbf{r}_1)$ being a proportionality factor and \mathbf{R} the tip-adatom lateral distance. Integrating over the surface coordinates, we find

$$\begin{aligned} I_1^\sigma &= 2e \int \frac{d\omega}{2\pi} \text{Re} \left\{ \sum_{\mathbf{k}_1} \int \int d\mathbf{r}_1 d\mathbf{r}'_1 e^{-i\mathbf{k}_1 \cdot \mathbf{r}_1} T^0(\mathbf{r}_1) \right. \\ &\quad \times e^{i\mathbf{k}_1 \cdot \mathbf{r}'_1} T^{0*}(\mathbf{r}'_1) [G_\sigma^r(\mathbf{R}, \mathbf{R}, \omega) g_{\mathbf{k}_1\sigma}^<(\omega) \\ &\quad \left. + G_\sigma^<(\mathbf{R}, \mathbf{R}, \omega) g_{\mathbf{k}_1\sigma}^a(\omega)] \right\}, \end{aligned} \quad (19)$$

and then over \mathbf{r}_1 and \mathbf{r}'_1 we arrive at

$$\begin{aligned} I_1^\sigma &= 2e \int \frac{d\omega}{2\pi} \text{Re} \left\{ \sum_{\mathbf{k}_1} |t_{13\mathbf{k}_1}^0|^2 [G_\sigma^r(\mathbf{R}, \mathbf{R}, \omega) g_{\mathbf{k}_1\sigma}^<(\omega) \right. \\ &\quad \left. + G_\sigma^<(\mathbf{R}, \mathbf{R}, \omega) g_{\mathbf{k}_1\sigma}^a(\omega)] \right\}, \end{aligned} \quad (20)$$

where $t_{13\mathbf{k}_1}^0$ is the Fourier transform of $T^0(\mathbf{r}_1)$. It is convenient to perform a Fourier transform on $G_\sigma^r(\mathbf{R}, \mathbf{R}, \omega)$ and $G_\sigma^<(\mathbf{R}, \mathbf{R}, \omega)$. This results in

$$\begin{aligned} I_1^\sigma &= 2e \int \frac{d\omega}{2\pi} \text{Re} \left\{ \sum_{\mathbf{k}_3, \mathbf{k}'_3} \sum_{\mathbf{k}_1} t_{13\mathbf{k}_1 \mathbf{k}_3}^0 t_{13\mathbf{k}_1 \mathbf{k}'_3}^* [G_{\mathbf{k}_3 \mathbf{k}'_3, \sigma}^r(\omega) g_{\mathbf{k}_1\sigma}^<(\omega) \right. \\ &\quad \left. + G_{\mathbf{k}_3 \mathbf{k}'_3, \sigma}^<(\omega) g_{\mathbf{k}_1\sigma}^a(\omega)] \right\}, \end{aligned} \quad (21)$$

where $t_{13k_1k_3} = t_{13k_1}^0 e^{i\mathbf{k}_3 \cdot \mathbf{R}}$.⁴⁰ In what follows, we assume $t_{13k_1}^0$ to be a constant, i.e., independent of \mathbf{k}_1 . As we have mentioned before, Eq. (21) could have been directly derived from the \mathbf{k} -space Hamiltonian $H_{13}^\sigma = \sum_{\mathbf{k}_1, \mathbf{k}_3} (t_{13} c_{\mathbf{k}_1\sigma}^\dagger c_{\mathbf{k}_3\sigma} + t_{13}^* c_{\mathbf{k}_3\sigma}^\dagger c_{\mathbf{k}_1\sigma})$, with $t_{13} = t_{13}^0 e^{i\mathbf{k}_3 \cdot \mathbf{R}}$. Observe that this is completely equivalent to the real-space Hamiltonian (8), with a point source, $T(\mathbf{r}_1, \mathbf{r}_3) = T^0(\mathbf{r}_1) \delta(\mathbf{r}_3 - \mathbf{R})$.

1. Spatially resolved transmission coefficient

Equation (20) can alternatively be written as

$$I_1^\sigma = e \int \frac{d\omega}{2\pi} [2\pi |t_{13}^0|^2 \rho_{1\sigma}(\omega)] i \{ G_\sigma^<(\mathbf{R}, \mathbf{R}, \omega) + f_1(\omega) [G_\sigma^r(\mathbf{R}, \mathbf{R}, \omega) - G_\sigma^a(\mathbf{R}, \mathbf{R}, \omega)] \}, \quad (22)$$

where $f_1(\omega)$ and $\rho_{1\sigma}(\omega) = \sum_{\mathbf{k}_1} \delta(\omega - \epsilon_{\mathbf{k}_1\sigma})$ are, respectively, the Fermi function and the density of states of the tip. If the tip is weakly coupled to the surface, we can calculate the current in the lowest order of $|t_{13}^0|^2$. This means that the Green functions $G_\sigma^r(\mathbf{R}, \mathbf{R}, \omega)$ and $G_\sigma^<(\mathbf{R}, \mathbf{R}, \omega)$ are assumed tip-decoupled, i.e., here, they are local Green functions for the subsystem adatom-surface only. In other words, the tip behaves as a probe, not affecting the local density of states. This allows us to write the current as

$$I_1^\sigma = e \int \frac{d\omega}{2\pi} [f_1(\omega) - f_3(\omega)] T_\sigma(\mathbf{R}, \omega), \quad (23)$$

where $f_3(\omega)$ is the Fermi function of the surface and the transmission coefficient is defined by

$$\begin{aligned} T_\sigma(\mathbf{R}, \omega) &= [2\pi |t_{13}^0|^2 \rho_{1\sigma}(\omega)] [-2\text{Im}\{G_\sigma^r(\mathbf{R}, \mathbf{R}, \omega)\}] \\ &= [2\pi |t_{13}^0|^2 \rho_{1\sigma}(\omega)] [2\pi \rho_\sigma(\mathbf{R}, \omega)], \end{aligned} \quad (24)$$

with $\rho_\sigma(\mathbf{R}, \omega) = -\frac{1}{\pi} \text{Im}\{G_\sigma^r(\mathbf{R}, \mathbf{R}, \omega)\}$ being the LDOS. In particular, in the absence of the adatom, the LDOS becomes the unperturbed surface density of states per spin, $\rho_3(\omega) = \sum_{\mathbf{k}_3} \delta(\omega - \epsilon_{\mathbf{k}_3}) = m/n\pi\hbar^2 = 1/2D = \rho_3^0$, with n being the two-dimensional (2D) electronic density and D the band half-width.

In this case and in the zero-temperature limit, the current reads

$$I_1^\sigma = e2\pi |t_{13}^0|^2 \rho_{1\sigma} \rho_3^0 eV, \quad (25)$$

where V is the bias voltage.

As we shall see in the next section, in the presence of the adatom, similarly to the case of an impurity on the surface of a metal, the LDOS shows Friedel oscillations; these affect the current that oscillates around the value given by Eq. (25).⁴¹

2. Calculating $\rho_\sigma(\mathbf{R}, \omega)$ for the subsystem adatom + surface

To determine the LDOS in the presence of the adatom, we calculate $G_\sigma^r(\mathbf{r}_3, \mathbf{r}'_3, \omega)$ by Fourier transforming $G_{\mathbf{k}_3\mathbf{k}'_3\sigma}^r(\omega)$,

$$G_\sigma^r(\mathbf{r}_3, \mathbf{r}'_3, \omega) = \sum_{\mathbf{k}_3\mathbf{k}'_3} e^{i\mathbf{k}_3 \cdot \mathbf{r}_3} e^{-i\mathbf{k}'_3 \cdot \mathbf{r}'_3} G_{\mathbf{k}_3\mathbf{k}'_3\sigma}^r(\omega), \quad (26)$$

assuming there is no tip-to-surface coupling, i.e., considering $H = H_2 + H_3 + H_{23}$. Following the procedure we described

in the previous section—equation of motion + analytical continuation—we obtain⁴³

$$\begin{aligned} G_{\mathbf{k}_3\mathbf{k}'_3\sigma}^r(\omega) &= \delta_{\mathbf{k}_3, \mathbf{k}'_3} g_{\mathbf{k}_3\sigma}^r(\omega) + \sum_{\mathbf{k}_2\mathbf{k}'_2} |t_{23}|^2 \\ &\quad \times g_{\mathbf{k}_3\sigma}^r(\omega) g_{\mathbf{k}'_2\sigma}^r(\omega) G_{\mathbf{k}_2\mathbf{k}'_2\sigma}^r(\omega), \end{aligned} \quad (27)$$

where $g_{\mathbf{k}_3\sigma}^r(\omega) = (\omega - \epsilon_{\mathbf{k}_3} + i\delta)^{-1}$ and $\sum_{\mathbf{k}_2\mathbf{k}'_2} G_{\mathbf{k}_2\mathbf{k}'_2\sigma}^r(\omega) \equiv G_{22\sigma}^r(\omega)$ is the adatom retarded Green function. Using Eq. (27) in Eq. (26), we find

$$\begin{aligned} G_\sigma^r(\mathbf{r}_3, \mathbf{r}'_3, \omega) &= \sum_{\mathbf{k}_3} \frac{e^{i\mathbf{k}_3 \cdot (\mathbf{r}_3 - \mathbf{r}'_3)}}{\omega - \epsilon_{\mathbf{k}_3} + i\delta} + |t_{23}|^2 \sum_{\mathbf{k}_3} \frac{e^{i\mathbf{k}_3 \cdot \mathbf{r}_3}}{\omega - \epsilon_{\mathbf{k}_3} + i\delta} \\ &\quad \times \sum_{\mathbf{k}'_3} \frac{e^{-i\mathbf{k}'_3 \cdot \mathbf{r}'_3}}{\omega - \epsilon_{\mathbf{k}'_3} + i\delta} G_{22\sigma}^r(\omega). \end{aligned} \quad (28)$$

Let

$$\sum_{\mathbf{k}_3} \frac{e^{i\mathbf{k}_3 \cdot \mathbf{r}_3}}{\omega - \epsilon_{\mathbf{k}_3} + i\delta} = \mathcal{R}(r_3, \omega) + iI(r_3, \omega), \quad (29)$$

where $\mathcal{R}(r_3, \omega)$ and $I(r_3, \omega)$ denote the corresponding real and imaginary parts given by

$$\mathcal{R}(r_3, \omega) = \rho_3^0 \int_{-1}^1 dx \frac{\frac{\omega}{D} - x}{(\frac{\omega}{D} - x)^2 + \delta^2} J_0(k_F r_3 \sqrt{1+x}) \quad (30)$$

and

$$I(r_3, \omega) = -\pi \rho_3^0 J_0 \left(k_F r_3 \sqrt{1 + \frac{\omega}{D}} \right), \quad (31)$$

with k_F being the Fermi wave number and $J_0(x)$ the Bessel function of the first kind. We can then write the LDOS as

$$\begin{aligned} \rho_\sigma(R, \omega) &= \rho_3^0 \left\{ 1 + \frac{\Gamma_3}{2} J_0^2 \left(k_F R \sqrt{1 + \frac{\omega}{D}} \right) \right. \\ &\quad \left. \times [(1 - q^2) \text{Im} G_{22\sigma}^r - 2q \text{Re} G_{22\sigma}^r(\omega)] \right\}, \end{aligned} \quad (32)$$

where $\Gamma_3 = 2\pi |t_{23}|^2 \rho_3^0$ and $q = \mathcal{R}(R, \omega)/I(R, \omega)$. In order to determine the transmission coefficient, we only have to calculate the adatom retarded Green function $G_{22\sigma}^r(\omega)$ obtained here using the Hubbard I approximation.³³ This approximation accounts for the Coulomb interaction and consists in factorizing the higher-order correlation functions appearing in the resulting equation of motion for $G_{22\sigma}^r(\omega)$. As a result we have³³

$$G_{22\sigma}^r(\omega) = \frac{1}{g_{2\sigma}^{r-1}(\omega) - \Sigma^r(\omega)}, \quad (33)$$

with

$$g_{2\sigma}^r(\omega) = \frac{\omega - \epsilon_\sigma - U(1 - n_{\bar{\sigma}})}{(\omega - \epsilon_\sigma)(\omega - \epsilon_\sigma - U)}, \quad (34)$$

where $\bar{\sigma} = -\sigma$, $n_{\bar{\sigma}}$ is the average occupation and Σ^r is the self-energy related to the coupling between the adatom and the host surface, $\Sigma^r = -\frac{i}{2}\Gamma_3$. Note that for the subsystem adatom + surface $n_{\bar{\sigma}} = n_\sigma$.

B. Resonant + nonresonant transport

In this section, we determine the *total* current—tip-to-adatom-to-host + tip-to-host—flowing in the system. In addition to the nonresonant (tip-to-host) current calculated in Sec. II A, here we consider the contribution from the direct tunneling of electrons between the tip and the adatom (resonant current). The corresponding additional term H_{12} is now taken into account and the Hamiltonian describing the system is given by Eq. (1). Note that for large enough R distances ($R \rightarrow \infty$) we regain the nonresonant case. Here, for convenience, we perform the calculation in \mathbf{k} space.

We model the tip-surface, tip-adatom, and adatom-surface couplings, respectively, by considering

$$t_{12} = t_{12}^0 e^{-R/R_0}, \quad (35)$$

$$t_{13} = t_{13}^0 e^{i\mathbf{k}_3 \cdot \mathbf{R}}, \quad (36)$$

$$t_{23} = t_{23}^0, \quad (37)$$

where t_{12}^0 , t_{13}^0 , and t_{23}^0 are (constant) phenomenological parameters and R_0 gives the exponential spatial decay for the coupling between the tip and the adatom as the tip moves away from it.

The current flowing into the tip or leaving it can be determined from Eq. (5). Calculating the commutator in this equation via Eqs. (1)–(3) we find

$$\begin{aligned} [H, N_1^\sigma] &= \sum_{j=2}^3 [H_{1j}, N_1^\sigma] \\ &= \sum_{j=2}^3 \sum_{\mathbf{k}_1, \mathbf{k}_j} (-t_{1j} c_{\mathbf{k}_1, \sigma}^\dagger c_{\mathbf{k}_j, \sigma} + t_{1j}^* c_{\mathbf{k}_j, \sigma}^\dagger c_{\mathbf{k}_1, \sigma}). \end{aligned} \quad (38)$$

Substituting this result into Eq. (5) we obtain

$$I_1^\sigma = -ie \sum_{j=2}^3 \sum_{\mathbf{k}_1, \mathbf{k}_j} [-t_{1j} \langle c_{\mathbf{k}_1, \sigma}^\dagger c_{\mathbf{k}_j, \sigma} \rangle + t_{1j}^* \langle c_{\mathbf{k}_j, \sigma}^\dagger c_{\mathbf{k}_1, \sigma} \rangle], \quad (39)$$

or in terms of the lesser Green function $G_{\mathbf{k}_j, \mathbf{k}_1, \sigma}^<(t, t)$,

$$I_1^\sigma = 2e \text{Re} \left[\sum_{j=2}^3 \sum_{\mathbf{k}_1, \mathbf{k}_j} t_{1j} G_{\mathbf{k}_j, \mathbf{k}_1, \sigma}^<(t, t) \right], \quad (40)$$

where $G_{\mathbf{k}_j, \mathbf{k}_1, \sigma}^<(t, t) = i \langle c_{\mathbf{k}_1, \sigma}^\dagger(t) c_{\mathbf{k}_j, \sigma}(t) \rangle$. Equation (40) is equivalent to Eq. (11) when $t_{12} = 0$. Now we must find an expression for the lesser Green function. Following the procedure described in Sec. II A, below we write down the equation of motion for the contour-ordered Green function $G_{\mathbf{k}_j, \mathbf{k}_1, \sigma}(\tau, \tau') = -i \langle T_C c_{\mathbf{k}_j, \sigma}(\tau) c_{\mathbf{k}_1, \sigma}^\dagger(\tau') \rangle$

$$\left(i \frac{\partial}{\partial \tau'} + \epsilon_{\mathbf{k}_1} \right) G_{\mathbf{k}_j, \mathbf{k}_1, \sigma}(\tau, \tau') = - \sum_{l=2}^3 \sum_{\mathbf{k}_l} t_{1l}^* G_{\mathbf{k}_j, \mathbf{k}_l, \sigma}(\tau, \tau'), \quad (41)$$

or in the integral form

$$G_{\mathbf{k}_j, \mathbf{k}_1, \sigma}(\tau, \tau') = \sum_{l=2}^3 \sum_{\mathbf{k}_l} t_{1l}^* \int_C d\tau_1 G_{\mathbf{k}_j, \mathbf{k}_l, \sigma}(\tau, \tau_1) g_{\mathbf{k}_1, \sigma}(\tau_1, \tau'), \quad (42)$$

where $g_{\mathbf{k}_1, \sigma}(\tau_1, \tau')$ is the tip free-electron Green function, and then analytically continue Eq. (42) to find $G_{\mathbf{k}_j, \mathbf{k}_1, \sigma}^<(t, t')$. Using this result in Eq. (40), we have

$$\begin{aligned} I_1^\sigma &= 2e \text{Re} \left\{ \sum_{j,l=2}^3 \sum_{\mathbf{k}_1, \mathbf{k}_j, \mathbf{k}_l} t_{1j} t_{1l}^* \int d\tau_1 [G_{\mathbf{k}_j, \mathbf{k}_l, \sigma}^r(t, \tau_1) g_{\mathbf{k}_1, \sigma}^<(t_1, t) \right. \\ &\quad \left. + G_{\mathbf{k}_j, \mathbf{k}_l, \sigma}^<(t, \tau_1) g_{\mathbf{k}_1, \sigma}^a(t_1, t) \right\}. \end{aligned} \quad (43)$$

Taking the Fourier transform of the above expression, we find

$$\begin{aligned} I_1^\sigma &= 2e \text{Re} \left\{ \int \frac{d\omega}{2\pi} \sum_{\mathbf{k}_1, \mathbf{k}_2, \mathbf{k}_3} t_{12} t_{12}^* [G_{\mathbf{k}_2, \mathbf{k}_2, \sigma}^r(\omega) g_{\mathbf{k}_1, \sigma}^<(\omega) \right. \\ &\quad + G_{\mathbf{k}_2, \mathbf{k}_2, \sigma}^<(\omega) g_{\mathbf{k}_1, \sigma}^a(\omega)] + \sum_{\mathbf{k}_1, \mathbf{k}_2, \mathbf{k}_3} t_{12} t_{13}^* [G_{\mathbf{k}_2, \mathbf{k}_3, \sigma}^r(\omega) g_{\mathbf{k}_1, \sigma}^<(\omega) \\ &\quad + G_{\mathbf{k}_2, \mathbf{k}_3, \sigma}^<(\omega) g_{\mathbf{k}_1, \sigma}^a(\omega)] + \sum_{\mathbf{k}_1, \mathbf{k}_3, \mathbf{k}_2} t_{13} t_{12}^* [G_{\mathbf{k}_2, \mathbf{k}_3, \sigma}^r(\omega) g_{\mathbf{k}_1, \sigma}^<(\omega) \\ &\quad + G_{\mathbf{k}_2, \mathbf{k}_3, \sigma}^<(\omega) g_{\mathbf{k}_1, \sigma}^a(\omega)] + \sum_{\mathbf{k}_1, \mathbf{k}_3, \mathbf{k}_3'} t_{13} t_{13}^* [G_{\mathbf{k}_3, \mathbf{k}_3', \sigma}^r(\omega) g_{\mathbf{k}_1, \sigma}^<(\omega) \\ &\quad \left. + G_{\mathbf{k}_3, \mathbf{k}_3', \sigma}^<(\omega) g_{\mathbf{k}_1, \sigma}^a(\omega) \right\}. \end{aligned} \quad (44)$$

Using Eqs. (35)–(37), we can rewrite Eq. (44) as

$$\begin{aligned} I_1^\sigma &= 2e \text{Re} \left\{ \int \frac{d\omega}{2\pi} |t_{12}^0|^2 e^{-2(R/R_0)} [G_{22\sigma}^r g_{1\sigma}^< + G_{22\sigma}^< g_{1\sigma}^a] \right. \\ &\quad + t_{12}^0 t_{13}^{0*} e^{-R/R_0} [G_{32\sigma}^r g_{1\sigma}^< + G_{32\sigma}^< g_{1\sigma}^a] \\ &\quad + t_{13}^0 t_{12}^{0*} e^{-R/R_0} [G_{23\sigma}^r g_{1\sigma}^< + G_{23\sigma}^< g_{1\sigma}^a] \\ &\quad \left. + |t_{13}^0|^2 [G_{33\sigma}^r g_{1\sigma}^< + G_{33\sigma}^< g_{1\sigma}^a] \right\}, \end{aligned} \quad (45)$$

where we have introduced the definitions

$$\begin{aligned} g_{1\sigma}^{<,a}(\omega) &= \sum_{\mathbf{k}_1} g_{\mathbf{k}_1, \sigma}^{<,a}(\omega), \\ G_{32\sigma}^{<,r} &= \sum_{\mathbf{k}_3, \mathbf{k}_2} e^{i\mathbf{k}_3 \cdot \mathbf{R}} G_{\mathbf{k}_3, \mathbf{k}_2, \sigma}^{<,r}, \\ G_{23\sigma}^{<,r} &= \sum_{\mathbf{k}_2, \mathbf{k}_3} e^{-i\mathbf{k}_3 \cdot \mathbf{R}} G_{\mathbf{k}_2, \mathbf{k}_3, \sigma}^{<,r}, \\ G_{33\sigma}^{<,r} &= \sum_{\mathbf{k}_3, \mathbf{k}_3'} e^{i(\mathbf{k}_3 - \mathbf{k}_3') \cdot \mathbf{R}} G_{\mathbf{k}_3, \mathbf{k}_3', \sigma}^{<,r}. \end{aligned} \quad (46)$$

Note that from Eq. (44) we regain Eq. (21) in the limit $t_{12}^0 = 0$, i.e., when the tip is far away from the adatom.

1. Matrix Green function formulation

We can see from Eq. (44) that $G_{\mathbf{k}_j, \mathbf{k}_1, \sigma}^<(t, t')$ is coupled to other Green functions. In order to find these Green functions, we have to apply the equation of motion technique to the corresponding contour-ordered Green function for each one of them and then perform an analytical continuation to obtain the

respective $G^{<,r}$. After a straightforward calculation we find

$$G_{\mathbf{k},\mathbf{k}_j,\sigma}(\tau,\tau') = \delta_{\mathbf{k},\mathbf{k}_j} g_{\mathbf{k}_j,\sigma}(\tau,\tau') + \sum_{l(l \neq j)} \sum_{\mathbf{k}_l} \int d\tilde{\tau} G_{\mathbf{k}_l,\mathbf{k}_j,\sigma}(\tau,\tilde{\tau}) t_{jl}^* g_{\mathbf{k}_j,\sigma}(\tilde{\tau},\tau'), \quad (47)$$

i.e., a system of coupled equations for the Green functions. Here, $g_{\mathbf{k}_j,\sigma}(\tau,\tau')$ is the free-electron Green function of the tip ($j = 1$), the adatom ($j = 2$), or the host surface ($j = 3$). These three Green functions $g_{\mathbf{k}_j,\sigma}(\tau,\tau')$ ($j = 1,2,3$) can be easily evaluated. Interestingly, by defining

$$G_{ij\sigma} = \sum_{\mathbf{k}_i,\mathbf{k}_j} G_{\mathbf{k}_i,\mathbf{k}_j,\sigma}, \quad (48)$$

$$G_{\underline{3}j\sigma} = \sum_{\mathbf{k}_3,\mathbf{k}_j} e^{i\mathbf{k}_3 \cdot \mathbf{R}} G_{\mathbf{k}_3,\mathbf{k}_j,\sigma}, \quad (49)$$

$$G_{j\underline{3}\sigma} = \sum_{\mathbf{k}_j,\mathbf{k}_3} e^{-i\mathbf{k}_3 \cdot \mathbf{R}} G_{\mathbf{k}_j,\mathbf{k}_3,\sigma}, \quad (50)$$

$$G_{\underline{3}\underline{3}\sigma} = \sum_{\mathbf{k}_3,\mathbf{k}'_3} e^{i(\mathbf{k}_3 - \mathbf{k}'_3) \cdot \mathbf{R}} G_{\mathbf{k}_3,\mathbf{k}'_3,\sigma}, \quad (51)$$

where the sum *is not* taken over the spin indices, we can write down a Dyson equation of the form

$$\mathbf{G}_\sigma(\tau,\tau') = \mathbf{g}_\sigma(\tau,\tau') + \int d\tau_1 \mathbf{G}_\sigma(\tau,\tau_1) \mathbf{\Sigma} \mathbf{g}_\sigma(\tau_1,\tau'), \quad (52)$$

with $\mathbf{G}_\sigma(\tau,\tau')$ being a matrix Green function whose elements are defined following Eqs. (48)–(51), i.e.,

$$\mathbf{G}_\sigma = \begin{pmatrix} G_{11\sigma} & G_{12\sigma} & G_{13\sigma} & G_{1\underline{3}\sigma} \\ G_{21\sigma} & G_{22\sigma} & G_{23\sigma} & G_{2\underline{3}\sigma} \\ G_{31\sigma} & G_{32\sigma} & G_{33\sigma} & G_{3\underline{3}\sigma} \\ G_{\underline{3}1\sigma} & G_{\underline{3}2\sigma} & G_{\underline{3}3\sigma} & G_{\underline{3}\underline{3}\sigma} \end{pmatrix}. \quad (53)$$

Additionally the self-energy is given by

$$\mathbf{\Sigma} = \begin{pmatrix} 0 & t_{12} & 0 & t_{13}^0 \\ t_{12}^* & 0 & t_{23} & 0 \\ 0 & t_{23}^* & 0 & 0 \\ t_{13}^{0*} & 0 & 0 & 0 \end{pmatrix} \quad (54)$$

and

$$\mathbf{g}_\sigma = \begin{pmatrix} g_{1\sigma} & 0 & 0 & 0 \\ 0 & g_{2\sigma} & 0 & 0 \\ 0 & 0 & g_{3\sigma} & g_{\underline{3}^*\sigma} \\ 0 & 0 & g_{\underline{3}\sigma} & g_{3\sigma} \end{pmatrix}, \quad (55)$$

with the matrix elements⁴⁴

$$g_{j\sigma}(\tau,\tau') = \sum_{\mathbf{k}_j} g_{\mathbf{k}_j,\sigma}(\tau,\tau'), \quad j = 1,2,3, \quad (56)$$

$$g_{\underline{3}\sigma}(\tau,\tau') = \sum_{\mathbf{k}_3} e^{i\mathbf{k}_3 \cdot \mathbf{R}} g_{\mathbf{k}_3,\sigma}(\tau,\tau'), \quad (57)$$

$$g_{\underline{3}^*\sigma}(\tau,\tau') = \sum_{\mathbf{k}_3} e^{-i\mathbf{k}_3 \cdot \mathbf{R}} g_{\mathbf{k}_3,\sigma}(\tau,\tau'). \quad (58)$$

Performing an analytic continuation in Eq. (52), we obtain the Dyson equation for the retarded Green function

$$\mathbf{G}_\sigma^r = [\mathbf{g}_\sigma^{r-1} - \mathbf{\Sigma}^r]^{-1} \quad (59)$$

and the Keldysh⁴⁵ equation

$$\mathbf{G}_\sigma^< = \mathbf{G}_\sigma^r \mathbf{g}_\sigma^{r-1} \mathbf{g}_\sigma^< \mathbf{g}_\sigma^{a-1} \mathbf{G}_\sigma^a, \quad (60)$$

where

$$\mathbf{g}_\sigma^{r,<} = \begin{pmatrix} g_{1\sigma}^{r,<} & 0 & 0 & 0 \\ 0 & g_{2\sigma}^{r,<} & 0 & 0 \\ 0 & 0 & g_{3\sigma}^{r,<} & g_{\underline{3}^*\sigma}^{r,<} \\ 0 & 0 & g_{\underline{3}\sigma}^{r,<} & g_{3\sigma}^{r,<} \end{pmatrix}. \quad (61)$$

The advanced Green function \mathbf{g}_σ^a is given by $\mathbf{g}_\sigma^a = [\mathbf{g}_\sigma^r]^*$. From Eqs. (59) and (60) we see that if \mathbf{g}_σ^r and $\mathbf{g}_\sigma^<$ are known we can determine immediately \mathbf{G}_σ^r and $\mathbf{G}_\sigma^<$, and so the spin-resolved current [see Eq. (45)]. The first nonzero (diagonal) elements ($g_{1\sigma}^<$ and $g_{1\sigma}^r$) in Eq. (61) are

$$g_{1\sigma}^<(\omega) = \sum_{\mathbf{k}_1} g_{\mathbf{k}_1,\sigma}^<(\omega) = \sum_{\mathbf{k}_1} 2\pi i f_1(\omega) \delta(\omega - \epsilon_{\mathbf{k}_1,\sigma}) = 2\pi i f_1(\omega) \rho_{1\sigma}(\omega), \quad (62)$$

$$g_{1\sigma}^r(\omega) = \sum_{\mathbf{k}_1} g_{\mathbf{k}_1,\sigma}^r(\omega) = \sum_{\mathbf{k}_1} \left[P \left(\frac{1}{\omega - \epsilon_{\mathbf{k}_1,\sigma}} \right) - i\pi \delta(\omega - \epsilon_{\mathbf{k}_1,\sigma}) \right] = \Lambda_1(\omega) - i\pi \rho_{1\sigma}(\omega), \quad (63)$$

where $f_1(\omega)$ is the tip Fermi distribution function, $\rho_{1\sigma}(\omega)$ is the tip density of states, and $\Lambda_1(\omega) = P \sum_{\mathbf{k}_1} \left(\frac{1}{\omega - \epsilon_{\mathbf{k}_1,\sigma}} \right)$, where P stands for the Cauchy principal value. The retarded adatom Green function $g_{2\sigma}^r(\omega)$ is given by Eq. (34). The lesser component can be calculated straightforwardly from the relation $g_{2\sigma}^<(\omega) = i n_\sigma A_\sigma(\omega)$, where

$$A_\sigma(\omega) = 2\pi(1 - n_\sigma) \delta(\omega - \epsilon_\sigma) + 2\pi n_\sigma \delta(\omega - \epsilon_\sigma - U), \quad (64)$$

and n_σ is the average spin-resolved occupation of the adatom. The third diagonal element of \mathbf{g}^r is given by

$$g_{3\sigma}^r(\omega) = \sum_{\mathbf{k}_3} g_{\mathbf{k}_3,\sigma}^r(\omega) = \sum_{\mathbf{k}_3} \left[P \left(\frac{1}{\omega - \epsilon_{\mathbf{k}_3}} \right) - i\pi \delta(\omega - \epsilon_{\mathbf{k}_3}) \right] = \Lambda_3(\omega) - i\pi \rho_3(\omega), \quad (65)$$

where $\rho_3(\omega) = \rho_3^0$ is the 2D density of states of the surface defined at the end of Sec. II A 1, and $\Lambda_3(\omega) = P \sum_{\mathbf{k}_3} \left(\frac{1}{\omega - \epsilon_{\mathbf{k}_3}} \right)$. For the corresponding lesser Green function, we find

$$g_{3\sigma}^<(\omega) = \sum_{\mathbf{k}_3} g_{\mathbf{k}_3,\sigma}^<(\omega) = 2\pi i f_3(\omega) \rho_3^0, \quad (66)$$

where $f_3(\omega)$ is the Fermi distribution function of the host surface.

Finally, we should calculate the off-diagonal elements of the matrix \mathbf{g}_σ . For the retarded Green function, we have

$$\begin{aligned} g_{3\sigma}^r(\omega) &= \sum_{\mathbf{k}_3} e^{i\mathbf{k}_3 \cdot \mathbf{R}} g_{\mathbf{k}_3\sigma}^r(\omega) \\ &= \rho_3^0 \int_{-1}^1 dx \frac{\frac{\omega}{D} - x}{\left(\frac{\omega}{D} - x\right)^2 + \delta^2} J_0(k_F R \sqrt{1+x}) \\ &\quad - i\pi \rho_3^0 J_0\left(k_F R \sqrt{1 + \frac{\omega}{D}}\right). \end{aligned} \quad (67)$$

For the lesser Green function we find

$$\begin{aligned} g_{3\sigma}^<(\omega) &= \sum_{\mathbf{k}_3} e^{i\mathbf{k}_3 \cdot \mathbf{R}} g_{\mathbf{k}_3\sigma}^<(\omega) \\ &= 2\pi i \rho_3^0 f_3(\omega) J_0\left(k_F R \sqrt{1 + \frac{\omega}{D}}\right). \end{aligned} \quad (68)$$

The Green functions $g_{3\sigma}^r$ and $g_{3\sigma}^<$ have exactly the same expressions as Eqs. (67) and (68), respectively.

C. Parameters and a summary of the numerical technique

Our main task is to determine the current from Eq. (45). To this end, we first calculate \mathbf{G}_σ^r and $\mathbf{G}_\sigma^<$ from Eqs. (59) and (60), respectively. Then we substitute the relevant matrix elements in Eq. (45). Note that in the presence of the Coulomb interaction, $g_{2\sigma}^r(\omega)$ and $g_{2\sigma}^<(\omega)$ depend on the adatom occupation n_σ ; so do \mathbf{G}_σ^r and $\mathbf{G}_\sigma^<$. This implies a self-consistent calculation, where n_σ is calculated iteratively via

$$n_\sigma = \int \frac{d\omega}{2\pi i} G_{22\sigma}^<(\omega). \quad (69)$$

As a matter of simplification we use the wide-band limit for the tip, so the density of states $\rho_{1\sigma}(\omega)$ is taken as constant ρ_1^0 (ρ_3^0 is already a constant), evaluated at the Fermi level. This is a good approximation when $eV, k_B T \ll D$, where D is the band half-width. The ferromagnetism of the tip is introduced via the density of states $\rho_{1\sigma} = \rho_1^0(1 \pm p)$, where p is the tip polarization and the $+$ and $-$ signs apply to spin up and down, respectively.³² Since the characteristic tunneling rate between the tip and the adatom is given by $\Gamma_{1\sigma} = 2\pi |t_{12}|^2 \rho_{1\sigma}$, we find $\Gamma_{1\sigma} = 2\pi |t_{12}|^2 \rho_1^0(1 \pm p)$, which is the standard phenomenology to account for the ferromagnetism of the electrode.⁴⁶ Analogously, the tunneling rate between the adatom and the host surface is $\Gamma_3 = 2\pi |t_{23}|^2 \rho_3^0 \equiv \Gamma_0$. In our calculation, we take $\Gamma_0 = 10 \mu\text{eV}$ as the energy scale. All the phenomenological parameters used in this paper are summarized in Table I.

TABLE I. Parameters used in the self-consistent calculation.

Parameter	Magnitude
Band half-width	$D = 1000\Gamma_0$
Adatom Charging Energy	$U = 30\Gamma_0$
Tip chemical potential	$\mu_1 = -eV/2$
Host chemical potential	$\mu_3 = eV/2$
System temperature	$k_B T = \Gamma_0$
Tip degree of polarization	$p = 0.4$
Decaying factor of t_{12}	$R_0 = 1/k_F$

Note that in Table I we define the origin of the energy scale such that $\mu_1 = \mu_3 = \epsilon_\sigma = 0$ at zero bias, i.e., the adatom energy level is aligned with the chemical potential of the leads in the absence of an applied bias. In order to be consistent with $\Gamma_0 = 2\pi |t_{23}|^2 \rho_3^0$ and the value of ρ_3^0 for typical semiconductors, e.g., GaAs, the factor t_{23} is taken as $\Gamma_0/\sqrt{10}$. We assume $t_{13}^0 = 0.01t_{23}^0$ and adopt values for t_{12}^0 and ρ_1^0 consistent with $\Gamma_0 = 2\pi |t_{12}^0|^2 \rho_1^0$. Note that for $eV = \pm 30\Gamma_0$ the adatom can be occupied by a single electron since ϵ_σ is within the conduction window (the energy range between μ_1 and μ_3) and $\epsilon_\sigma + U$ is without this range. On the other hand, for $eV = \pm 150\Gamma_0$, the dot can be doubly occupied since both ϵ_σ and $\epsilon_\sigma + U$ lie inside the conduction window. The parameter R_0 controls how fast the coupling t_{12} decays in space when the tip moves away from the adatom. We take it equal to k_F^{-1} . Hereafter k_F^{-1} will be used as a length scale.

III. RESULTS

A. Single Occupancy

Figure 2(a) shows the adatom occupations against the lateral distance R between the tip and the adatom. We study both

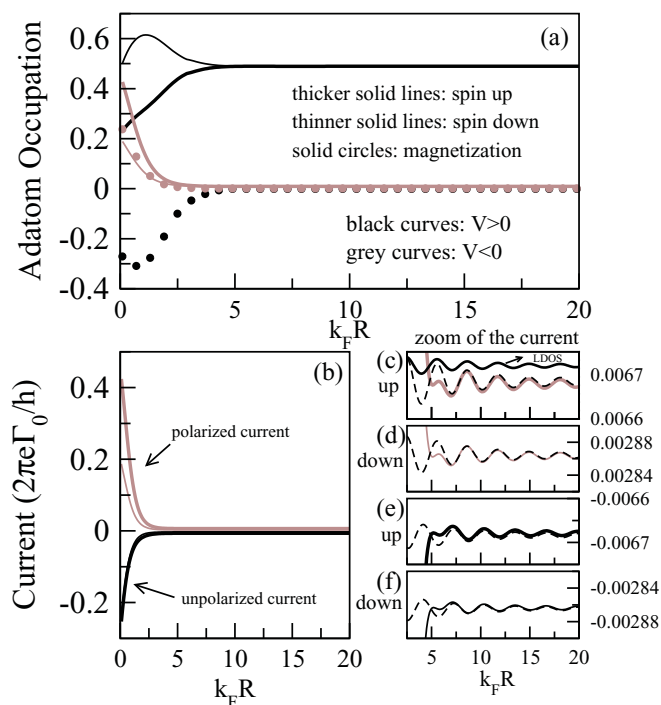


FIG. 2. (Color online) (a) Adatom occupations and magnetization and (b) spin-resolved currents against the tip position R . Both negative (grey line) and positive (black line) bias voltages are shown. For $|eV| = 30\Gamma_0$, only single occupancy (i.e., $\epsilon_\sigma + U$ is much higher than the emitter chemical potential) is allowed. This results in the spin-diode effect: for negative bias voltages the current is polarized for all R values while for positive biases the current polarization goes to zero when the tip is close to the adatom. Panel (c)–(f): zoom of each spin component of the current in the range where the tip-adatom coupling is negligible. In (c) we also show the LDOS (thick solid black line) evaluated at ϵ_F . The LDOS displays Friedel oscillations which show up in the polarized current. The vertical scale for the LDOS is not shown.

forward ($eV = +30\Gamma_0$) and reverse ($eV = -30\Gamma_0$) bias cases. As mentioned before, for $|eV| = 30\Gamma_0$ the adatom cannot be doubly occupied since $\epsilon_\sigma + U$ lies above the Fermi energy of the source. For $V > 0$, the host NM surface is the source and the FM tip is the drain of electrons, i.e., the electrons flow from the NM surface (+ adatom) to the tip. For $V < 0$, we have the opposite.

For $R = 0$ and $V > 0$ (black lines), we find $n_\uparrow < n_\downarrow$. This is reasonable since spin-up electrons tunnel from the adatom into the tip much more easily than the spin-down ones, due to the larger spin-up density of states, $\rho_{1\sigma} = \rho_1^0(1 \pm p)$ ($p > 0$), which implies $\Gamma_{1\uparrow} > \Gamma_{1\downarrow}$, and gives rise to a larger spin-down population. For a constant bias voltage, as the tip moves away from the adatom [see Eq. (35)], the tunneling rates $\Gamma_{1\uparrow}$ and $\Gamma_{1\downarrow}$ decrease, but the incoming rates Γ_3 stay the same, this results in an increase of both n_\uparrow and n_\downarrow . We note, however, that the spin-down population (thinner solid black line) increases more quickly than the spin-up one (thicker solid black line), this is because $\Gamma_{1\downarrow} < \Gamma_{1\uparrow}$, see Fig. 2(a). As the n_\uparrow adatom population increases with R , the n_\downarrow one tends to be more blocked due to the spin-dependent Coulomb blockade. The interplay between the Coulomb blockade and the decrease of the tunneling rates $\Gamma_{1\uparrow,\downarrow}$ makes n_\downarrow reach a maximum, subsequently decreasing to attain the limit $n_\uparrow = n_\downarrow = 0.5$ for large enough R 's.

In contrast, for $eV = -30\Gamma_0$, $n_\uparrow > n_\downarrow$ for small R values, see in Fig. 2(a) the solid thicker and thinner grey lines. This is a consequence of $\Gamma_{1\uparrow} > \Gamma_{1\downarrow}$, which means that more spins up tunnel to the adatom. Besides, the outgoing rates Γ_3 (the same for the up and down components) is smaller than $\Gamma_{1\uparrow}$, which results in a larger spin-up accumulation in the adatom. As the tip moves away from the adatom, $\Gamma_{1\uparrow}$ and $\Gamma_{1\downarrow}$ go to zero exponentially and the populations n_\uparrow and n_\downarrow are completely drained out into the host surface, thus resulting in an empty adatom.

The magnetization $m = n_\uparrow - n_\downarrow$ is also shown in Fig. 2(a) (solid circles). Observe that for small R the adatom is spin-down polarized for $V > 0$ and spin-up polarized for $V < 0$. As R increases, m tends to zero for both positive and negative bias voltages. However, m tends to zero much slower for $V > 0$ than for $V < 0$, a consequence of the interplay between the Coulomb interaction (spin-Coulomb blockade) and the tunneling rates $\Gamma_{1\sigma}$, that change with the tip position as it moves away from the adatom.

In Fig. 2(b), we present the spin-resolved currents for both $eV = \pm 30\Gamma_0$. The spin-diode effect²⁴ can be clearly seen for small values of R . While for $V > 0$ (black lines) we find $I_\uparrow \approx I_\downarrow$ for small R values, for $V < 0$ (grey lines) we observe $I_\uparrow > I_\downarrow$. This shows that the current polarization can be controlled via both the bias sign and the tip position. In the case of $V < 0$, we have $\Gamma_{1\uparrow} > \Gamma_3 > \Gamma_{1\downarrow}$, i.e., the spin-up population is greater than the spin-down one, $m > 0$. As a consequence, in the absence of the Coulomb interaction in the adatom, $I_\uparrow > I_\downarrow$ (the case $U = 0$ resembles the curves in the double-occupancy regime ($eV \gg U$), see Fig. 3). In the presence of U , I_\downarrow is suppressed since n_\downarrow tends to be more blocked than n_\uparrow [see Fig. 2(a)], which results in an enhancement in the difference between I_\uparrow and I_\downarrow . For $V > 0$, the magnetization changes sign, $m < 0$, now the spin-up population tends to be more blocked and I_\uparrow is more strongly suppressed compared to I_\downarrow , interestingly, attaining values close to I_\downarrow . The amplification

of I_\uparrow compared to I_\downarrow for $V < 0$, when the tip is closer to the adatom, does not occur in the double occupancy regime ($eV = \pm 150\Gamma_0$) as we will see in the next section.

In Figs. 2(c)–2(f), we show the current for a range of R in which only the direct tip-host tunneling (nonresonant transport) is relevant. Note that I_\uparrow and I_\downarrow tend to distinct plateaus for large enough R 's. These plateaus correspond to the background current between the tip and the host surface, given approximately by Eq. (25). By comparison with Eq. (45) we plot in dashed line the current obtained via Eq. (23). In the large- R limit, we expect an agreement between both equations, since Eq. (23) was derived in the case of negligible tunneling between the tip and the adatom (see the solid black and grey lines). The minor difference between the two results is due to Eq. (23) having been obtained in the limit of small tip-surface coupling parameter t_{13}^0 .⁴¹ The LDOS evaluated at the Fermi level, $\rho_\sigma(R, 0)$ [Eq. (32)], is also shown in Fig. 2(c); it oscillates around the unperturbed surface density of states ρ_3^0 . Friedel-like oscillations are seen for both spin components, thus reflecting the oscillations in the LDOS due to the scattering center (adatom). Note that Friedel oscillations have been seen experimentally in a variety of systems.^{26–28,30}

B. Double Occupancy

Figure 3 shows the spin-resolved (a) adatom occupations and (b) currents in the double-occupancy regime, i.e., when the bias voltage is large enough ($|eV| = 150\Gamma_0$) to allow for two electrons of opposite spins in the adatom at the same

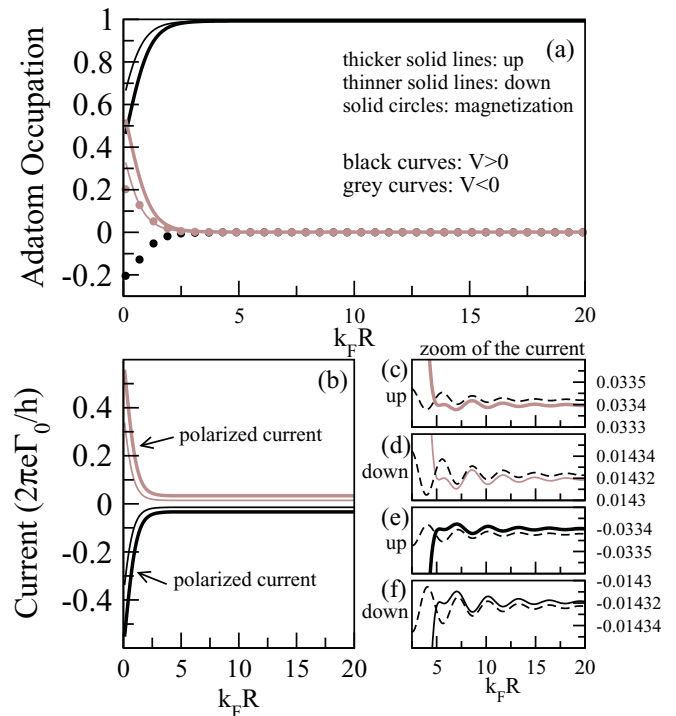


FIG. 3. (Color online) Similar to Fig. 2 except that $|eV| = 150\Gamma_0$ here. This bias allows double occupancy of the adatom. In this regime no spin-diode effect is observed. The spin polarized currents are the same (in modulus) for both positive and negative eV . For forward voltages (black curves), the adatom becomes doubly occupied for large R values ($n_\uparrow + n_\downarrow = 2$).

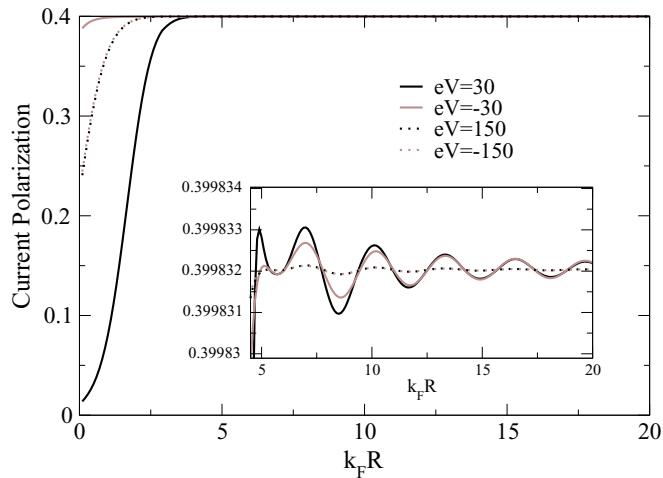


FIG. 4. (Color online) Current polarization against R . In the single-occupancy regime ($eV = \pm 30\Gamma_0$) the current polarization is approximately constant for reverse bias ($V < 0$) and goes down to zero for direct bias ($V > 0$) when the tip is close to the adatom. In the case of double occupancy ($eV = \pm 150\Gamma_0$), the current polarization is suppressed for small R , however, it does not vanish; notice that the dotted lines lie essentially on top of each other. For large R 's, the polarization for both single and double occupancies tends to a plateau and presents tiny oscillations (inset).

time. For $V > 0$ (black lines), as R increases, both n_\uparrow and n_\downarrow populations tend to unity and the adatom becomes doubly occupied ($n_\uparrow + n_\downarrow = 2$). This is so because electrons can jump into the adatom but cannot leave it for large R values. In contrast, for $V < 0$ both n_\uparrow and n_\downarrow vanish as R increases because the electron source (tip) decouples from the adatom. Observe also that the magnetization m is enhanced as R tends to zero and switches sign depending on whether V is positive or negative.

The current in the double-occupancy regime [see Fig. 3(b)] has a similar behavior for both positive and negative biases. Note that $|I_\uparrow| > |I_\downarrow|$ for $V > 0$ and $V < 0$, in contrast to the single-occupancy regime where we find $I_\uparrow \approx I_\downarrow$ for $V > 0$ [see Fig. 2(b)]; hence, no spin-diode effect is observed here. In Figs. 3(c)–3(f), we show a zoom of the spin-resolved currents in the range of negligible tip-adatom coupling. As before, we observe Friedel oscillations, which reflect the disturbance in the LDOS due to the localized impurity (adatom). The dashed black lines in Figs. 3(c)–3(f) show the current obtained via Eq. (23).

C. Current polarization

Figure 4 displays the current polarization, $\wp = (I_\uparrow^\uparrow - I_\uparrow^\downarrow)/(I_\uparrow^\uparrow + I_\uparrow^\downarrow)$, as a function of R . Both single ($|eV| = 30\Gamma_0$) and double ($|eV| = 150\Gamma_0$) occupancies are shown. For $eV = -30\Gamma_0$ (solid grey line) the polarization is approximately constant $\sim 40\%$ while for $eV = +30\Gamma_0$ (solid black line) it is strongly suppressed when the tip is close to the adatom. This is a type of spatially resolved spin-diode²⁴ that allows a polarized current to flow only for reverse bias. In the case of double occupancy, though, both positive and negative biases present a similar behavior with a 40% current polarization away from the adatom and a slight suppression as the tip moves closer to it. This 40% current polarization for large R 's in the single- and double-occupancy cases follows straightforwardly by calculating \wp using the spin-resolved nonresonant currents in Eq. (25). The inset shows a blow up of the current polarization and also reveals Friedel oscillations.

IV. CONCLUSION

We have studied spin-polarized quantum transport in a system composed of an FM-STM tip coupled to a NM host surface with a single adsorbed atom. Due to Coulomb interaction in the adatom, the system can operate as a spin-diode when the tip is nearby the adatom. In the singly occupied case and direct bias ($V > 0$), the current polarization can vary from zero up to 40% depending on the tip position. For reverse bias, though, the polarization is pinned close to 40% for all tip positions. In the double occupancy regime, the current polarization is the same for both forward and reverse biases, with a slight suppression as the tip moves closer to the adatom. Additionally, the adatom magnetization can be tuned by varying the tip position and its sign can switch depending on the bias. Finally, we have also found spin-resolved Friedel oscillations in the current as the tip moves laterally away from the adatom, thus reflecting the oscillations in the surface LDOS induced by the adatom acting like an effective impurity.

ACKNOWLEDGMENTS

Two of the authors, J.C.E. and P.H.P., acknowledge useful discussions with D. Loss. The authors acknowledge financial support from CNPq, CAPES, FAPEMIG, and FAPESP.

¹E. Meyer, H. J. Hug, and R. Bennewitz, *Scanning Probe Microscopy: The Laboratory on a Tip* (Springer-Verlag, Berlin, 2004).

²*Atomic and Nanometer Scale Modification of Materials*, edited by Ph. Avouris (Kluwer, Dordrecht, 1993).

³C. Julian Chen, *Introduction to Scanning Tunneling Microscopy* (Oxford University Press, Oxford, 1993).

⁴M. F. Crommie, C. P. Lutz, and D. M. Eigler, *Science* **262**, 218 (1993).

⁵H. C. Manoharan, C. P. Lutz, and D. M. Eigler, *Nature (London)* **403**, 512 (2000).

⁶D. Awschalom and M. E. Flatté, *Nat. Phys.* **3**, 153 (2007).

⁷D. Serrate, P. Ferriani, Y. Yoshida, S.-W. Hla, M. Menzel, K. von Bergmann, S. Heinze, A. Kubetzka, and R. Wiesendanger, *Nat. Nanotechnology* **5**, 350 (2010).

⁸C. Çelebi, J. K. Garleff, A. Yu. Silov, A. M. Yakunin, P. M. Koenraad, W. Van Roy, J.-M. Tang, and M. E. Flatté, *Phys. Rev. Lett.* **104**, 086404 (2010).

- ⁹D. Kitchen, A. Richardella, J.-M. Tang, M. E. Flatté, and A. Yazdani, *Nature (London)* **442**, 436 (2006).
- ¹⁰A. R. Smith, *J. Scanning Probe Microscopy* **1**, 3 (2006).
- ¹¹H. Oka, P. A. Ignatiev, S. Wedekind, G. Rodary, L. Niebergall, V. S. Stepanyuk, D. Sander, and J. Kirschner, *Science* **327**, 843 (2010).
- ¹²N. O. Dantas, E. S. F. Neto, R. S. Silva, D. R. Jesus, and F. Pelegrini, *Appl. Phys. Lett.* **93**, 193115 (2008).
- ¹³M. Kamenetska, M. Koentopp, A. C. Whalley, Y. S. Park, M. L. Steigerwald, C. Nuckolls, M. S. Hybertsen, and L. Venkataraman, *Phys. Rev. Lett.* **102**, 126803 (2009).
- ¹⁴F. Mohn, J. Repp, L. Gross, G. Meyer, M. S. Dyer, and M. Persson, *Phys. Rev. Lett.* **105**, 266102 (2010).
- ¹⁵G. Schull, T. Frederiksen, M. Brandbyge, and R. Berndt, *Phys. Rev. Lett.* **103**, 206803 (2009).
- ¹⁶R. A. Žak, B. Röthlisberger, S. Chesi, and D. Loss, *Riv. Nuovo Cimento* **33**, 345 (2010).
- ¹⁷S. Loth, M. Etzkorn, C. P. Lutz, D. M. Eigler, and A. J. Heinrich, *Science* **329**, 1628 (2010).
- ¹⁸A. A. Khajetoorians and A. Kubetzka, *Nat. Nanotechnology* **5**, 830 (2010).
- ¹⁹M. Gierling, P. Schneeweiss, G. Visanescu, P. Federsel, M. Häffner, D. P. Kern, T. E. Judd, A. Günther, and J. Fortágh, *Nat. Nanotechnology* **6**, 446 (2011).
- ²⁰V. Madhavan, W. Chen, T. Jamneala, M. F. Crommie, and N. S. Wingreen, *Science* **280**, 567 (1998).
- ²¹Nikolaus Knorr, M. Alexander Schneider, Lars Diekhöner, Peter Wahl, and Klaus Kern, *Phys. Rev. Lett.* **88**, 096804 (2002).
- ²²K. R. Patton, S. Kettemann, A. Zhuravlev, and A. Lichtenstein, *Phys. Rev. B* **76**, 100408(R) (2007).
- ²³A. C. Seridonio, F. M. Souza, J. Del Nero, and I. A. Shelykh, *Physica E* **41**, 1611 (2009).
- ²⁴F. M. Souza, J. C. Egues, and A. P. Jauho, *Phys. Rev. B* **75**, 165303 (2007).
- ²⁵C. A. Merchant and N. Marković, *Phys. Rev. Lett.* **100**, 156601 (2008).
- ²⁶Ph. Avouris, I.-W. Lyo, R. E. Walkup, and Y. Hasegawa, *J. Vac. Sci. Technol. B* **12**, 1447 (1994).
- ²⁷K. Kanisawa, M. J. Butcher, H. Yamaguchi, and Y. Hirayama, *Phys. Rev. Lett.* **86**, 3384 (2001).
- ²⁸I. Affleck, L. Borda, H. Saleur, *Phys. Rev. B* **77**, 180404(R) (2008).
- ²⁹A. C. Seridonio, F. S. Orahcio, F. M. Souza, and M. S. Figueira, e-print [arXiv:1106.2853v2](https://arxiv.org/abs/1106.2853v2) (unpublished).
- ³⁰S. L. Kawahara, J. Lagoute, V. Repain, C. Chacon, Y. Girard, J. Klein, and S. Rousset, *Phys. Rev. B* **82**, 020406(R) (2010).
- ³¹E. C. Stoner, *Proc. R. Soc. London A* **169**, 339 (1939).
- ³²F. M. Souza, J. C. Egues, and A. P. Jauho, *Braz. J. Phys.* **34**, 565 (2004).
- ³³H. Haug and A. P. Jauho, *Quantum Kinetics in Transport and Optics of Semiconductors*, Springer series in Solid-State Sciences 123 (Springer, New York, 1996).
- ³⁴J. Rammer and H. Smith, *Rev. Mod. Phys.* **58**, 323 (1986).
- ³⁵H. Bruus and K. Flensberg, *Many-body Quantum Theory in Condensed Matter Physics: an Introduction* (Oxford University Press, New York, 2004).
- ³⁶D. C. Langreth, *Linear and Nonlinear Electron Transport in Solids*, edited by J. T. Devreese (Plenum, New York, 1976).
- ³⁷G. A. Fiete and E. J. Heller, *Rev. Mod. Phys.* **75**, 933 (2003).
- ³⁸J. Tersoff and D. R. Hamann, *Phys. Rev. B* **31**, 805 (1985).
- ³⁹M. Plihal and J. W. Gadzuk, *Phys. Rev. B* **63**, 085404 (2001).
- ⁴⁰The proportionality $t_{13} \propto e^{ik_3 \cdot \mathbf{R}}$ was originally discussed in Ref. 38 and further used in Ref. 39.
- ⁴¹The current plateau given by Eq. (25) is valid only for small tip-surface coupling parameter t_{13}^0 . If we calculate the Green functions $G_\sigma^r(\mathbf{R}, \mathbf{R}, \omega)$ and $G_\sigma^<(\mathbf{R}, \mathbf{R}, \omega)$ accounting for the tip, a small correction arises and Eq. (25) becomes $I_1^\sigma = G_\sigma V$, where $G_\sigma = \frac{\varepsilon}{\hbar} \frac{4r}{(1+r)^2}$, with $r = \pi^2 |t_{13}^0|^2 \rho_{1\sigma} \rho_3$. This feature was pointed out in Ref. 42. Note that we can regain Eq. (25) (now divided by \hbar) by expanding the above expression for the current up to second order in the coupling parameter.
- ⁴²E. Vernek, N. Sandler, and S. E. Ulloa, *Phys. Rev. B* **80**, 041302(R) (2009).
- ⁴³To obtain the equation of motion for $G_{\mathbf{k}_3 \mathbf{k}_3, \sigma}^r(t - t')$, we first perform a time derivative with respect to t ; the resulting equation couples $G_{\mathbf{k}_3 \mathbf{k}_3, \sigma}^r(t - t')$ to $G_{\mathbf{k}_2 \mathbf{k}_3, \sigma}^r(t - t')$. By deriving this second Green function now with respect to t' , we find an equation that relates $G_{\mathbf{k}_3 \mathbf{k}_3, \sigma}^r(t - t')$ to the adatom Green function $G_{\mathbf{k}_2 \mathbf{k}_2, \sigma}^r(t - t')$.
- ⁴⁴Note that Eq. (27) and the LDOS, given by Eq. (32), can be derived from the present matrix formulation by taking $t_{12} = 0$ and $t_{13} = 0$ in the self-energy matrix [see Eq. (54)].
- ⁴⁵Q. F. Sun, J. Wang, and H. Guo, *Phys. Rev. B* **71**, 165310 (2005).
- ⁴⁶W. Rudziński and J. Barnás, *Phys. Rev. B* **64**, 085318 (2001).