Four-terminal quantum dot as an efficient rectifier of heat and charge currents

Karol I. Wysokiński D

Institute of Physics, M. Curie-Skłodowska University, pl. M. Curie-Skłodowskiej 1, 20-031 Lublin, Poland

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We propose an efficient method of heat rectification in a simple system consisting of a quantum dot asymmetrically coupled to four mutually perpendicular electrodes. In such a device, the perpendicular charge and heat currents appear in response to the voltage bias or temperature difference between one pair of electrodes. Even though both longitudinal (along the bias) and perpendicular (to the bias) currents are rectified under appropriate conditions, the rectification factor is typically much bigger for the latter currents. This is true for heat and charge flow. The perpendicular currents are predicted to exist in linear as well as nonlinear transport regimes and require broken mirror symmetry but not time reversal symmetry. The linear effect exists only in geometry which breaks two inversion symmetries along two pairs of electrically coupled terminals. The proposed system is attainable within current technology and provides a novel platform of simultaneous heat and charge management at the nanoscale.

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I. INTRODUCTION

A rectifier is a device in which the magnitude of the charge or heat current depends on the sign of the electric or thermal bias. Efficient heat current rectifiers are among the most desirable devices at the nanoscale as they allow for heat management in miniaturized electronic devices. Various geometries and devices have been proposed to reach the goal. An early proposal of nanoscale heat rectification has appeared in the context of molecular electronics [1] and is still debated in the literature [2,3].

In recent years an increased interest is observed in theoretical analysis and experimental studies of thermal diodes [4]. Besides molecules including those with negative U centers [5], many systems with tuned quantum properties have been proposed as possible rectifiers [6]. These include inter alia recently analysed two terminal junctions with bath particles obeying different particle exchange statistics [7] and the use of quantum entanglement as a possible tools to enhance rectification properties [8]. Also various novel solid state systems and materials have been put forward. These include heterostructures, functionally graded or phase changing normal materials, superconductors, etc. as reviewed recently [9,10].

In the context of this paper, the rectifying devices based on quantum dots are of special interest [11–24]. They consist of a single or more quantum dots coupled to two external reservoirs. Probably the first experimental demonstration of rectifying properties of a two terminal quantum dot is that in Ref. [25] inferred from the asymmetric line shape of the thermopower. Rectification of both charge and heat currents in the Coulomb blockade regime in the system with two quantum dots has been studied in Ref. [21]. In Ref. [24], the authors concentrate on heat rectification through quantum dots in the Coulomb blockade regime using master equation approach. They considered two-terminal and four-terminal devices. In the latter case, two coupled quantum dots form a main nanoscopic element in which each of the individual dots is contacted by two separate terminals. Our four-terminal system is different, as it relies on two pairs of electrodes coupled to a single quantum dot. To the best of our knowledge, it has not been studied before in the context of rectification. The geometry allows for a new functionality which is the rectification of both longitudinal and perpendicular currents. This property might be of importance in nanosystems with many terminals.

Quantum dots play an important role in novel electronic devices like single electron transistors [26], heat nanoengines [27], and many more, including building blocks of quantum computers [28,29]. Quantum dots with large charging energy U, coupled to external metallic leads behave like magnetic impurities in noble metals [30,31] and at low temperature show Kondo effect [32,33].

The device we are proposing consists of a single quantum dot tunnel coupled to four normal electrodes in a cross geometry as shown in Fig. 1. The application of the voltage or thermal bias along LR electrodes (i.e., x direction) results in a simultaneous flow of (longitudinal) charge and heat currents between L and R electrodes and also between U and D electrodes (perpendicular currents) if the system breaks mirror symmetry(-ies). A device with nonsymmetrical couplings works as an efficient rectifier. The rectification is observed for both currents and directions. If the interaction U of electrons is non-negligible, as it is usually the case in small structures, and if working temperature is low enough, the device allows to study the effect of Kondo correlations on the longitudinal and perpendicular currents.

Four terminal nanojunctions with similar geometry have been studied previously. General symmetry properties of phase coherent transport in the presence of magnetic flux were derived in Ref. [34]. The same geometry was theoretically analysed in the context of perpendicular currents and resistances [35,36] taking into account polarization of electrons and strong spin-orbit interaction in the central region. Wei



FIG. 1. The four terminal quantum dot geometry used in this work. The couplings Γ^{λ} between the central dot and the terminals $(\lambda = L, R, U, D)$ take on arbitrary values. The external leads can differ by the chemical potentials μ_{λ} and/or temperatures T_{λ} . Occasionally we denote the direction along *L* and *R* terminals as the *x* direction and that along *U* and *D* as the *y* direction.

et al. studied planar four terminal system [37] and found the nonlinear Hall effect induced by the dipole of Berry curvature. The hybrid structure consisting of a quantum dot contacted by tunnel barriers to four electrodes including two superconducting and two normal leads, allows [38] the control of supercurrent flowing between a pair of superconducting electrodes by the bias voltage applied to normal electrodes. The cross geometry of the four terminal nanojunction containing an asymmetric prismlike scatterer has been studied experimentally [39]. Even though our system is different from the experimental one it features similar antisymmetric dependence of the four terminal resistance on the current as discussed below.

We shall analyze two possible boundary conditions: open with the floating U and D electrodes, and closed with flow of the perpendicular current. There is no perpendicular voltage [40] under open boundary conditions. However, the currents perpendicular to the direction of the bias exist under the closed boundary conditions. These conditions are necessary but not sufficient for the observation of the perpendicular currents. Breaking of single or two mirror symmetries is needed. The system with broken symmetries shows rectification properties. The rectification efficiency may attain its maximal possible value equal unity. Tuning the system to such hot spots allows perfect rectification of charge or heat. We hope this property can be verified experimentally as the fabrication of the proposed devices is possible by the present-day technology.

The organization of the paper is as follows. In Sec. II, we describe the studied device and its modeling. Two possible boundary conditions are discussed in Secs. II A and II B. The results of numerical calculations presented in Section (III) are followed by the concluding section, Sec. IV. Some detailed calculations of currents flowing in the system and the on-dot Green function are relegated to Appendices A and B.

II. THE GEOMETRY, CURRENTS AND BOUNDARY CONDITIONS

The time reversal symmetric nanostructure consisting of a quantum dot tunnel-coupled to four normal leads is illustrated in Fig. 1. The quantum dot is understood here as the small grain with quantized spectrum, which can be modelled by a single level. Electrodes or terminals are macroscopic objects characterized by temperature $T_{\lambda} = T + \Delta T_{\lambda}$. They may be electrically biased (eV_{λ}) with chemical potential $\mu_{\lambda} = \mu + eV_{\lambda}$, where μ is an equilibrium value of the chemical potential and *T* the equilibrium temperature common to all electrodes. We assume $\mu = 0$.

The Hamiltonian describing the system under consideration is a standard single-impurity Anderson model

$$H = \sum_{\lambda k\sigma} \varepsilon_{\lambda k} n_{\lambda k\sigma} + \sum_{\sigma} \varepsilon_{d\sigma} n_{\sigma} + U_{d} n_{\uparrow} n_{\downarrow} + \sum_{\lambda k\sigma} (V_{\lambda k\sigma} c^{\dagger}_{\lambda k\sigma} d_{\sigma} + V^{*}_{\lambda k\sigma} d^{\dagger}_{\sigma} c_{\lambda k\sigma}), \qquad (1)$$

where $n_{\lambda k\sigma} = c_{\lambda k\sigma}^{\dagger} c_{\lambda k\sigma}$ and $n_{\sigma} = d_{\sigma}^{\dagger} d_{\sigma}$ denote particle number operators for the leads and the dot, respectively. The operators $c_{\lambda k\sigma}^{\dagger}(d_{\sigma}^{\dagger})$ create electrons in respective states $\lambda k\sigma$ (σ) in the lead λ (on the dot). The energies of the leads are measured from their chemical potentials μ_{λ} , $\varepsilon_{\lambda k} = \varepsilon_{0\lambda k} - \mu_{\lambda}$, with the dependence of $\varepsilon_{0\lambda k}$ on λ allowing for a different spectrum in each of the leads. U_d denotes the energy cost of placing two electrons on a quantum dot. In this work, we neglect spin dependence of the on dot energy $\varepsilon_{d\sigma} = \varepsilon_d$ and electron hoping amplitudes $V_{\alpha k\sigma} = V_{\alpha k}$. These approximations are relaxed in hybrid system with normal and ferromagnetic electrodes or in the presence of magnetic field. To study the transport properties, we employ the nonequilibrium Greens function technique. We express the currents flowing in the system by the equilibrium Fermi distribution functions with parameters adequate to corresponding terminals, the effective couplings between the dot and terminals and the retarded dot Green function (for details see Appendixes A and B).

A. Absence of Hall voltage in systems with open boundary conditions

We start by assuming the bias *B* (this denotes either voltage *V* or temperature ΔT) between *L* and *R* electrodes. For open boundary conditions, chemical potentials μ_U and μ_D are obtained by requiring that the currents in those electrodes vanish: $I_U = 0$ and $I_D = 0$. We calculate these currents by using the general equation (A2) and express the result in terms of the integrals defined in equation (A4). Neglecting the spin dependence, it is easy to show that under arbitrary voltage bias V_{LR} the vanishing of charge currents in the *U* and *D* electrodes requires

$$F_D - F_U = 0 = \int \frac{dE}{2\pi} (f_D(E) - f_U(E)) N(E).$$
(2)

In the above formula, the density of states, defined in Eq. (A6), is the nonequilibrium one. Due to the fact that N(E) is a strictly positive function of energy, the vanishing of the integral (2) requires equality of the equilibrium Fermi distribution functions at the considered terminals: $f_D(E) = f_U(E)$. For

equal temperatures ($T_D = T_U$) the only solution is $\mu_D = \mu_U$ and no voltage appears along *UD* direction, regardless the parameters of the model. This is valid for arbitrary voltages and temperatures, i.e., in the linear and nonlinear regimes. Thus under open boundary conditions there is no "Hall voltage" in the system and thus no true Hall effect. This, however, does not preclude flow of the currents along *UD* direction.

B. The perpendicular currents in the four terminal QD with closed boundary conditions

The situation is completely different if one applies closed boundary conditions and allows for a current flow along y direction. We bias a system along the x direction with $V = V_R - V_L$ or $\Delta T = T_R - T_L$ and calculate the current flow along the y, i.e., UD direction. It has to be noted that the voltage bias induces both, heat J^Q and charge I currents along LR and UD and the same is true for the temperature bias. The currents are calculated as $I^{LR} = (I^L - I^R)/2$ and $I^{UD} = (I^U - I^D)/2$ for charge and $J_Q^{LR} = (J_Q^L - J_Q^R)/2$, $J_Q^{UD} = (J_Q^U - J_Q^D)/2$ for a heat flow.

The necessary condition for the existence of currents along unbiased *UD* direction is broken mirror symmetry between *U* and *D* electrodes. This is realized by assuming different couplings to the dot. For a system with $\Gamma^U = \Gamma^D$, the perpendicular currents vanish, $I^{UD} = 0$ ($J_Q^{UD} = 0$).

This general statement can be easily proved. Calculating both, the heat and charge currents flowing along the UDdirection in the same way as explained in Sec. II A, one finds that they can be generally expressed as

$$i^{UD} = (\Gamma^U - \Gamma^D) f_i(\{\Gamma^\lambda\}, \{V_\lambda\}, \{\Delta T_\lambda\})$$
(3)

and thus vanish for $\Gamma^D = \Gamma^U$. Here f_i is the numerically evaluated function for a particular current, and i^{UD} is used to denote either I^{UD} or J_Q^{UD} . It is also of interest to note that the currents are expected to change sign if the difference $\Gamma^U - \Gamma^D$ changes sign. It has to be stressed, however, that this is not the universal feature as the function $f_i(\{\Gamma^\lambda\}, \{V_\lambda\}, \{\Delta T_\lambda\})$ which is a combination of the appropriate integrals defined in (A4) or (A5) disturbs the said proportionality depending on the parameters. For a particular example, where one observes no sign change of the currents $I^{UD}(V)$ calculated for two sets of Γ^D and Γ^U despite sign change of $\Gamma^D - \Gamma^U$ see Sec. (III B).

In the linear response, the conditions for the existence of the perpendicular currents are stronger and require breaking of two mirror symmetries. This is easily seen assuming over a range of voltages isothermal conditions ($T_{\lambda} = T$ for all λ) and the linear response regime (small voltage $V = V_L - V_R$). For symmetric distribution of voltages $V_{L/R} = \pm V/2$ at the corresponding terminals, one expands the Fermi functions

$$f_{\lambda}(E) = f_0(E) + f'_0(E)(-eV_{\lambda}), \tag{4}$$

and using Eq. (A2) gets the perpendicular charge current

$$I^{UD} = \frac{4e^2}{h} \frac{(\Gamma^U - \Gamma^D)[\Gamma^L - \Gamma^R]}{\sum_{\lambda} \Gamma^{\lambda}} F_0' V, \qquad (5)$$

where $F'_0 = \int \frac{dE}{2\pi} (-\frac{\partial f_0(E)}{\partial E}) N(E)$. Both currents vanish for $\Gamma^L = \Gamma^R$ and/or $\Gamma^U = \Gamma^D$. This condition is relaxed in the

nonlinear regime and breaking the mirror symmetry along UD is enough to get the perpendicular currents.

III. THE RESULTS

The asymmetry of the couplings plays an important role and in most cases we shall characterize it by a single parameter α , which defines anisotropy of our system. It may take arbitrary positive value but we shall study a few representative values only. In most studied cases, we assume simple asymmetry with $\Gamma^R = \Gamma^D = \alpha \Gamma_0$ and $\Gamma^L = \Gamma^U = \Gamma_0$, where Γ_0 is our energy unit. The calculations for the couplings less symmetric give similar values of the currents. We stick to the simplest choices, which allow to study the role which symmetries play in our model. We also assume Boltzmann constant, Planck constant, and the electron charge as $k_B = \hbar = e = 1$. Thus energy E and other parameters like *T*, *V* and ε_d , are all measured in units of Γ_0 .

A. The currents

Contrary to the linear regime in which existence of perpendicular currents require breaking of two mirror symmetries, in the nonlinear regime both charge I^{ij} and heat J_Q^{ij} currents flow between ij = LR as well as ij = UD electrodes if only a single mirror symmetry is broken along UD direction. This is illustrated in Fig. 2 for voltage bias and three values $\Gamma^D = 0.5, 1.5, 2$ with all other couplings equal to $\Gamma_0 = 1$. In accord with Eq. (5) for $\Gamma^L = \Gamma^R$, the linear contributions vanish and the perpendicular currents are (at least) quadratic functions of voltage for small V. Simultaneously the longitudinal currents (both I^{LR} and J_Q^{LR}) are linear functions of V for $V \to 0$ with well visible departures from linearity at larger voltages [see insets in Fig. 2(a) and 2(b)]. This agrees with general analytical results of Sec. II B. Thermally induced currents (not shown) exhibit the same behavior, namely, the currents perpendicular to the bias are quadratic functions of ΔT for small ΔT . These currents do not appear in the linear order if the system breaks the single mirror symmetry only. In full analogy to the voltage bias, the longitudinal thermally induced currents are linear functions of ΔT for $\Delta T \rightarrow 0$, with departures from linearity at elevated ΔT values.

For the particular set of parameters, the *UD* heat current is positive for $\Gamma^D > 1$ and negative for $\Gamma^D < 1$. This is true independently of the bias as visible from panel (b) in Fig. 2. Similar symmetry is valid for heat currents along *LR*, which for $\Gamma^D > 1$ are of opposite sign to the currents corresponding to $\Gamma^D < 1$. The flow towards *U* is either blocked or facilitated.

The response of the system to applied bias quite generally depends on its symmetry and the set of parameters. To illustrate this in Fig. 3, we show thermally induced charge currents in panel (a) and heat currents induced by the voltage in panel (b). Panel (a) shows the currents in system at temperature T = 2 with the quantum dot tuned to $\varepsilon_d = -9$, and for two values of anisotropy $\alpha = 0.5$, 1.5 and U = 12. The currents flowing along the *UD* are about an order of magnitude smaller then those along *LR* direction. However, one notices that the charge perpendicular current for $\alpha = 0.5$ vanishes for $\Delta T/2 \approx 0.4$ but is finite for $\Delta T/2 \approx -0.4$. This is a bias for which the rectification factor is maximal (=1) as



FIG. 2. Charge current I^{UD} induced by voltage between *LR* terminals is a symmetric (and roughly quadratic) function of *V* as visible in (a). The same is true for heat current which is also symmetric (and quadratic) function of voltage (b). Insets show the corresponding currents along *LR* direction. These currents are linear functions at small voltages with nonlinear behavior observed at high bias. Moreover, the charge currents along *RL* are very weakly affected by the ratio Γ^D/Γ^U . Other parameters read T = 1, $\varepsilon_d = -4$, U = 12, and $\Gamma^R = \Gamma^L = \Gamma^U = \Gamma_0$. All energies are measured in units of Γ_0 .

we shall see in the following section. In panel (b), we show perpendicular heat currents versus the voltage. Important fact to note is the nonmonotonous dependence of currents on the bias and the points were they vanish. These special points are marked by asterisks.

B. Rectification

If mirror symmetries along *LR* and *UD* directions are broken both charge and heat currents in two pairs of electrodes depend on the sign of bias B = V or $B = \Delta T$, i.e. we find $I^{LR}(B) \neq I^{LR}(-B)$ and $I^{UD}(B) \neq I^{UD}(-B)$. The same is true for heat currents $J_Q^{LR}(B)$ and $J_Q^{UD}(B)$. To quantify the rectification efficiency one introduces special parameter called rectification coefficient. One possible definition is

$$R^{ij} = \frac{|(|I^{ij}(B)| - |I^{ij}(-B)|)|}{[|I^{ij}(B)| + |I^{ij}(-B)|]}.$$
(6)

With this definition the rectification factor ranges from 0 to 1. The former means no rectification, while the latter perfect rectification. Similar rectification coefficients are defined for



FIG. 3. In (a), we show thermally induced longitudinal charge currents I^{LR} (solid lines) and perpendicular currents I^{UD} (lines with symbols) for two values of asymmetry $\alpha = 0.5$ and $\alpha = 1.5$ as functions of the temperature difference. In (b), the voltage dependent perpendicular heat currents J_Q^{UD} are shown together with points marked with * where the currents vanish for one sign of voltages and take on nonzero values for opposite sign. In both panels, U = 12 in units of Γ_0 .

heat currents $J^{Q}(B)$ and we denote them by R_{Q}^{ij} in the following.

Perfect rectification is expected in cases when the current vanishes for one sign of the bias and attains finite value for the other. The goal is to tune the parameters of the system to such "hot spots." In the panel (b) of Fig. 3 such hot spots have been marked by asterisks. The rectification coefficient takes on the maximal possible value at those points. In general it is difficult to tune the system and find vanishing of charge currents. However, panel (a) of this figure shows points in which perpendicular charge currents vanish for specific values of ΔT . For $\alpha = 0.5$, the I^{UD} current vanishes for $\Delta T/2$ close to 0.4, while for $\alpha = 1.5$, the perpendicular charge currents are relatively large for opposite sign of temperature bias.

The rectification coefficients for thermally induced charge and heat currents flowing along *LR* and *UD* directions are plotted in Fig. 4. Panel (a) shows R^{ij} for charge currents, while panel (b) R_Q^{ij} for heat currents. The rectification of both currents flowing along the bias direction (I^{LR} and J_Q^{LR}) is relatively small of order of a few percent. For the parameters of the model presented in the figure, the coefficient R^{UD} is of the same order or slightly higher then R^{LR} . Its maximal value is



FIG. 4. Rectification ratios R^{ij} of charge (a) and heat R_Q^{ij} (b) currents as a function of $\Delta T/2$ for a few values of α , voltage V = 0, temperature T = 1, $\varepsilon_d = -4$, and U = 12.

about 10%. However, the heat rectification coefficient $R_Q^{UD} \gg R_Q^{LR}$ and reaches values higher then 50%. It monotonously increases with ΔT .

Temperature plays an important role in our interacting system, as the Kondo effect sets in at low temperature. To see this, we show in Fig. 5 results obtained for the same system as in Fig. 4 with $\alpha = 0.5$ and for two vastly different temperatures. We plot rectification factors R^{ij} (main panel) and the charge currents I^{ij} (inset) versus voltage V for two values of temperature. For lower temperature T = 0.01, the



FIG. 5. Rectification ratios R^{ij} and currents I^{ij} (inset) as a function of voltage for two temperatures T = 1 (solid lines) and 0.01 (lines with symbols). Other parameters $\varepsilon_d = -4$ and U = 12.

Kondo resonances are expected to appear in the density of states.

The signature of the Kondo effect is visible in the inset of Fig. (5) as a higher value of the current at a given voltage at low T compared to the same current at higher temperature. Concomitantly the slopes $(\partial I/\partial V)$ of the curves $I^{UD}(V)$ and $I^{LR}(V)$ calculated for low temperature (T = 0.01 curves with symbols, when a Kondo effect is expected to appear) are higher then the slopes of the same currents calculated at high temperatures (T = 1 thin solid lines when there is no Kondo effect) leading to higher values of the low temperature conductances. The effect is best visible close to V = 0. Also at higher voltages the different values of currents show the influence of the Kondo correlations despite the fact that in the strongly nonlinear regime one calculates the current by integrating the density of states over large energy window and this makes the relative contribution of the Kondo resonances smaller. We note in passing that the Kondo temperature estimated from the Haldane's formulas (A7) for the parameters used in Fig. 5 is found $T_K \approx 0.7$. Another way to see the Kondo effect on the current directly is to plot the latter as a function of temperature for a constant value of the voltage. As a result of the calculations (not shown) one gets the curve which at temperature T well below the Kondo temperature T_K shows constant value of the current. With increasing temperature toward T_K the current decreases in the expected manner.

Strong nonlinearities at elevated voltages combined with well visible asymmetry of I^{UD} current (at low T = 0.01) for voltage around $|V^*| \approx 1.2$ result in $|I^{UD}(V^*)| = |I^{UD}(-V^*)|$ and vanishing of $R^{UD}(V^*)$ as well as its nonmonotonous dependence on voltage. At high temperature T = 1, there is no Kondo effect and one obtains continuous increase of rectification with V. The rectification of longitudinal current R^{LR} is affected by the Kondo effect only quantitatively. For the particular set of parameters, its low temperature value is roughly doubled with respect to higher temperature but remains low, at the level of a few percent.

The general observation is that typically the rectification factors for longitudinal currents are small of the order of a few percent. This agrees with previous systematic calculations of the rectification factor in Ref. [24] in a two terminal single level quantum dot. The rectification factors were small of order of a few percent like those for longitudinal currents in the present work. In our four terminal geometry, the rectification factor for the perpendicular currents is usually much higher and may be as large 100%.

In view of the observation [24] that the rectification coefficient for a two level quantum dot attains large value in the two terminal system it would be of interest to extend our four terminal model to two level quantum dot and to study how the longitudinal and perpendicular currents and their rectification factors are affected. This will be studied in a future work.

C. Four terminal resistance in the nonlinear regime

For many terminal nanojunctions with currents I^{ij} and voltages V^{kl} measured between various pairs of electrodes one defines [34], the four terminal resistances $R_{ij,kl} = \frac{V^{kl}}{I^{ij}}$. The voltages are applied between terminals k and l and currents measured between terminals i and j. Here we shall



FIG. 6. Broken mirror symmetry along vertical direction only $\Gamma^R = \Gamma^L = \Gamma^U = 1$ and $\Gamma^D = 0.5$, 1.5, and 2 results in the four terminal resistivity antisymmetric with respect to the current flowing along horizontal direction. This resembles experimental result [39] found in the nonlinear transport regime of nanojunction with artificial asymmetric scatterer.

calculate resistances for a special couplings which break a mirror symmetry along y direction only. If we assume couplings to L, R, and U terminals equal to Γ_0 and only the coupling to $\Gamma^D = (0.5, 1.5, 2)\Gamma_0$ different from others, the resulting symmetry of the model resembles that of the nanojunction earlier studied experimentally [39]. These authors considered a four terminal structure with asymmetric triangular prism-like scatterer placed in its center. The scatterer effectively blocked the flow of charge from/to one of the terminals. In our case it is the coupling Γ^D which effectively blocks (if <1) the current from/to this terminal. It should be recalled that in the linear transport regime the resistances $R_{ii,kl}$ do not depend on the current. In the nonlinear regime, the perpendicular current I^{UD} depends on the longitudinal current I^{LR} and so does the resistance $R_{UD,LR} = \frac{V^{LR}}{I^{UD}} = R_{UD,LR}(I^{LR})$. Due to the vanishing of the perpendicular currents at some voltages in our set-up, the four terminal resistance $R_{UD,LR}$ is not well defined at those points. That is why we show in Fig. 6 the inverse resistance $R_{UD,LR}^{-1} = \frac{I^{UD}}{V^{LR}}$ as a function of the current $I = I^{LR}$. It is seen that the resistance obeys a symmetry $R_{UD,LR}(I) = -R_{UD,LR}(-I)$. Interestingly, similar symmetry of the four terminal resistance has been earlier observed [39] in the nonlinear ballistic transport in the (already mentioned) four terminal microjunction with triangular asymmetric (prismlike) scatterer. The absence of voltage V^{UD} between U and D electrodes in our model and its presence in experimental setup is a main reason of the perfect symmetry $R_{UD,LR}(I) = -R_{UD,LR}(-I)$ in our case and only an approximate one in the nanojunction [39].

Inverse resistances $R_{UD,LR}^{-1}$ calculated in the Kondo regime characterized by the parameters $\varepsilon_d = -4$, U = 12, and T = 0.01 display the same perfect antisymmetric dependence (not shown) on the longitudinal current I^{LR} . Needless to say that this is true for the mirror symmetry broken along UD direction in a similar way as in experiment [39].

D. Gate voltage dependence and the effect of interactions

In the nonlinear regime, the density of states (DOS) defined in (A6) and entering formula (A4) is known to depend



FIG. 7. The dependence of the longitudinal I^{LR} (main panel) and transverse I^{UD} (inset) currents on the detuning $\delta = \varepsilon_d + U/2$ (can be changed by the gate voltage) for system with $\Gamma^L = \Gamma^U = \Gamma_0$ and $\Gamma^R = \Gamma^D = 1.5\Gamma_0$ (i.e., $\alpha = 1.5$), U = 16, voltage bias V = 4and for two temperatures T = 0.01 and 0.5. Signatures of the Kondo effect visible as non smooth dependence of the currents on δ are observed for T = 0.01.

crucially on the interactions U between carriers, voltages V_{λ} and temperatures T_{λ} of the leads. We limit the studies of this section to isothermal condition when all leads have the same temperature $T_{\lambda} = T$. If temperature is low enough and the on-dot level ε_d is slightly below the chemical potential(s), the Kondo peak(s) appears in the density of states of the interacting quantum dot. With chemical potentials $\mu = 0$ at electrodes U and D, $\mu_{L/R} = \mu \pm eV/2$ at the left/right electrode one expects three Kondo peaks pinned to the chemical potentials of the electrodes. They are visible in Fig. 9 shown in Appendix B. Various curves in the figure correspond to different values of $\Gamma^R = \Gamma^D = \alpha \Gamma_0$ couplings with $\Gamma^L = \Gamma^U = \Gamma_0$. For ε_d outside the Kondo regime DOS around E = 0 changes smoothly with α and voltage V (not shown).

In Fig. 7, we show the currents I^{LR} (main panel) and I^{UD} (inset) as a function of dot energy ε_d at temperatures below (T = 0.01) and above (T = 0.5) the Kondo temperature. The dot energy can be easily changed by gate voltage. The system breaks mirror symmetry with respect to both LR and UD directions as we have assumed $\Gamma^L = \Gamma^U = \Gamma_0$ and $\Gamma^R = \Gamma^D =$ $1.5\Gamma_0$. At lower temperature one observes signatures of the Kondo effect. These are peaks for those gate voltages for which one expects the Kondo resonances in the density of states (cf. Fig. 9). The effect is rather small due to the fact that we integrate the density of states over a range (-V/2, V/2)around E = 0. In this energy window, there are three Kondo peaks including one well pronounced and voltage independent at E = 0. It is bounded with the chemical potential $\mu = 0$ of both U and D electrodes. Interestingly, the effect of Kondo correlations is more pronounced in the perpendicular current I^{UD} as compared to longitudinal one I^{LR} .

In the upper panel of color coded Fig. 8, we show the dependence of longitudinal I^{LR} charge current on $\delta = \varepsilon_d + U/2$ and voltage V. The lower panel shows similar dependence for the perpendicular charge current I^{UD} . Both currents are calculated for temperature in the Kondo regime (T = 0.01) and for U = 12. Note the same scale for both panels, however, with perpendicular current multiplied by the factor of 10.



FIG. 8. The maps show the charge currents on the plane (δ, V) . Upper panel illustrates longitudinal current $I^{LR}(\varepsilon_d, V)$, while lower panel the perpendicular current $I^{UD}(\delta, V)$. We assumed here U = 12and temperature T = 0 and 0.01 in energy units.



FIG. 9. The evolution of the on-dot density of states plotted as a function of energy with changing the parameter α describing the couplings between the dot and right (*R*) and down (*D*) terminals with $\Gamma^R = \Gamma^D = \alpha \Gamma_0$, $\Gamma^L = \Gamma^U = \Gamma_0$. The parameters take on the following values: the on-dot energy $\varepsilon_d = -5.0$, the source-drain voltage V = 2, temperature T = 0.01 and the interaction U = 12. Note that the region of width V around the chemical potential $\mu = 0$ is important as it contributes to the currents in particular terminals.

TABLE I. Necessary conditions for the rectification of the currents along *LR* and *UD* directions if the bias (voltage or temperature) is applied between the *L* and *R* electrodes. The couplings Γ^{λ} which are not mentioned can take on arbitrary values.

	linear regime	nonlinear regime
$I^{LR}; J^{LR}_O$	$\Gamma^L \neq \Gamma^R$	$\Gamma^L \neq \Gamma^R$
$I^{UD}; ilde{J}_Q^{UD}$	$\Gamma^L \neq \Gamma^R, \Gamma^U \neq \Gamma^D$	$\Gamma^U \neq \Gamma^D$

IV. SUMMARY AND CONCLUSION

We have studied the nonlinear transport characteristics of a system consisting of strongly interacting quantum dot coupled to two pairs of normal leads arranged in a cross geometry. The voltage or thermal bias applied to one pair of the leads induces the (heat and charge) current flow between both pairs of terminals, provided the quantum dot is nonsymmetrically coupled (cf. Table I). Different couplings between the leads and the dot break mirror symmetries of the device. In the linear regime observation of two mutually perpendicular (charge and heat) currents in response to the appropriate bias requires breaking of both symmetries. Beyond linear regime, breaking the mirror symmetry along UD is enough. All currents are rectified in a system breaking both mirror symmetries, with the rectification factors of perpendicular currents typically much bigger then that of longitudinal currents. Interestingly the rectification factor for heat typically exceed that for charge.

If only the symmetry between vertical leads (perpendicular to those biased) is broken the resulting nonlinear four terminal resistance $R_{ij,kl} = \frac{V^{kl}}{I^{ij}}$ is an antisymmetric function of the longitudinal current $I = I^{LR}$, i.e., $R_{UD,LR}(I) = -R_{UD,LR}(-I)$. This is in qualitative agreement with experimental data [39] on nonlinear transport in four terminal microjunction with asymmetric prismlike scatterer. The notable difference is that in our system the antisymmetry is exact while in experiment it is approximate. It is important to recall that in the linear regime $R_{ij,kl}$ does not depend on the current $I = I^{ij}$.

We want also to underline that the analyzed device with a cross geometry enables the study of the Kondo effect on both longitudinal and perpendicular currents. It turns out that the Kondo resonance appearing in the density of states at low temperature affects the longitudinal currents in the four terminal system to lesser extend then in the corresponding two terminal quantum dot geometry. Additionally in our geometry the signatures of the Kondo effect in the perpendicular current are more pronounced then in the longitudinal one.

Our results demonstrate a new route to achieve the efficient rectification of longitudinal and perpendicular (heat and charge) currents. The proposed system is attainable within current technology and provides a novel platform of simultaneous heat and charge management at the nanoscale.

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APPENDIX A: THE CURRENT IN THE SYSTEM

Calculating the currents we follow the general definitions relying on equation of motion of the number and energy operators [41,42]. The heat and charge currents are expressed by the lesser and greater Green functions, which have to be calculated for the fully interacting system. This is a difficult task, especially for the strongly nonlinear regime, we are interested in here. The required lesser Green function is calculated in a standard way [43–45] on the time contour. It turns out that the calculations can be simplified in the so called wide band limit. In this limit, one assumes that the couplings between the dot and the leads $\Gamma^{\lambda}_{\sigma}(E) = 2\pi \sum_{k} |V_{\lambda k\sigma}|^2 \delta(E - \varepsilon_{\lambda k}) = \Gamma^{\lambda}_{\sigma}$ do not depend on energy. In this limit, one finds the exact relation, first derived in Ref. [46]

$$\begin{aligned} \langle d_{\sigma}^{\dagger} d_{\sigma} \rangle &= -i \int \frac{dE}{2\pi} G_{\sigma}^{<}(E) \\ &= i \int \frac{dE}{2\pi} \frac{\sum_{\lambda} \Gamma_{\sigma}^{\lambda} f_{\lambda}(E)}{\sum_{\lambda} \Gamma_{\sigma}^{\lambda}} [G_{\sigma}^{r}(E) - G_{\sigma}^{a}(E)], \, (A1) \end{aligned}$$

which allows to write the charge and heat currents flowing out of the λ electrode as (for details, see Refs. [45,46])

A 15

$$I_{\lambda} = \frac{2e}{\hbar} \int \frac{dE}{2\pi} \sum_{\sigma} \Gamma_{\sigma}^{\lambda}$$

$$\times \frac{\sum_{\lambda'} \Gamma_{\sigma}^{\lambda'}(f_{\lambda'}(E) - f_{\lambda}(E))}{\sum_{\lambda'} \Gamma_{\sigma}^{\lambda'}} \text{Im} G_{\sigma}^{r}(E), \quad (A2)$$

$$J_{\lambda} = \frac{2e}{\hbar} \int \frac{dE}{2\pi} \sum_{\sigma} \Gamma_{\sigma}^{\lambda}(E)(E - \mu_{\lambda})$$

$$\times \frac{\sum_{\lambda'} \Gamma_{\sigma}^{\lambda'}(f_{\lambda'}(E) - f_{\lambda}(E))}{\sum_{\lambda'} \Gamma_{\sigma}^{\lambda'}} \text{Im} G_{\sigma}^{r}(E).$$
(A3)

These expressions can be used for calculating the currents in an arbitrary system consisting of the central dot and several terminals under appropriate boundary conditions. Note the Fermi-Dirac distribution function $f_{\lambda}(E)$) depends on the electrode λ via its chemical potential $\mu_{\lambda} = \mu + eV_{\lambda}$ or applied voltage V_{λ} and temperature T_{λ} . The common value of the chemical potential and temperature of the system in equilibrium is denoted μ , T. For generality we keep here the spin dependence of adequate parameters.

It is convenient to express the currents in terms of auxiliary integral

$$F_{\lambda\sigma} = \int \frac{dE}{2\pi} f_{\lambda}(E) N_{\sigma}(E), \qquad (A4)$$

$$F^{\mathcal{Q}}_{\lambda\sigma} = \int \frac{dE}{2\pi} (E - \mu_{\lambda}) f_{\lambda}(E) N_{\sigma}(E), \qquad (A5)$$

respectively. In the above formulas,

$$N_{\sigma}(E) = -\frac{1}{\pi} \operatorname{Im} G_{\sigma}^{r}(E, \{V_{\lambda}\})$$
(A6)

denotes density of states (DOS) on the dot for spin σ electrons. In general, this quantity depends on the voltages $\{V_{\lambda}\}$ and temperatures T_{λ} of all terminals.

As our theory is valid also in the Kondo regime we finish this Appendix with quoting the Haldane's formula [47] for the Kondo temperature expected in the system

$$T_K = 0.5\sqrt{U\Gamma_N} \exp\left(\frac{\pi\varepsilon_d(\varepsilon_d + U)}{2\Gamma_N U}\right),\tag{A7}$$

with $\Gamma_N = \sum_{\lambda} \Gamma^{\lambda}$.

APPENDIX B: THE DOT GREEN FUNCTION

For completeness, we recall the formulas for the on-dot Green function which qualitatively correctly describe the Kondo effect. One uses standard equation of motion technique [44–46] and finds the dot Green function

$$G_{\sigma}^{r}(E) = \langle \langle d_{\sigma} | d_{\sigma}^{\dagger} \rangle \rangle_{E}^{r} = \frac{1 + I_{d}(E)[\langle n_{\bar{\sigma}} \rangle + b_{1\bar{\sigma}} - b_{2\bar{\sigma}}]}{E - \varepsilon_{\sigma} - \Sigma_{0\sigma} + \Sigma_{t}(E)},$$
(B1)

where

$$\Sigma_t(E) = I_d(E) \Big[\Sigma_{1\sigma}^T + \Sigma_{2\sigma}^T - (b_{1\bar{\sigma}} - b_{2\bar{\sigma}}) \Sigma_{0\sigma} \Big], \qquad (B2)$$

$$I_d(E) = \frac{U}{E - \varepsilon_{\sigma} - U - \Sigma_{0\sigma} - \Sigma_{\sigma}^{(1)} - \Sigma_{\sigma}^{(2)}}.$$
 (B3)

The various pieces of the self-energy are supplemented by the inverse life-times $i\gamma_{\bar{\sigma}_1/2}$ of the single particle $\bar{\sigma}$, respectively two-particle 2 state and read

$$b_{1\bar{\sigma}}(E) = \int \frac{d\varepsilon}{2\pi} \frac{\sum_{\lambda} \Gamma_{\bar{\sigma}}^{\lambda} f_{\lambda}(\varepsilon) \langle \langle d_{\bar{\sigma}} | d_{\bar{\sigma}}^{\dagger} \rangle \rangle_{\varepsilon}^{a}}{E - \varepsilon - \varepsilon_{1} + i \tilde{\gamma}_{1}^{\bar{\sigma}}}, \qquad (B4)$$

$$b_{2\bar{\sigma}}(E) = \int \frac{d\varepsilon}{2\pi} \frac{\sum_{\lambda} \Gamma_{\bar{\sigma}}^{\lambda} f_{\lambda}(\varepsilon) \langle \langle d_{\bar{\sigma}} | d_{\bar{\sigma}}^{\dagger} \rangle \rangle_{\varepsilon}^{a}}{E + \varepsilon - \varepsilon_{2} + i\tilde{\gamma}_{2}}, \qquad (B5)$$

$$\Sigma_{1\sigma}^{T}(E) = \int \frac{d\varepsilon}{2\pi} \frac{\sum_{\lambda} \Gamma_{\tilde{\sigma}}^{\lambda} f_{\lambda}(\varepsilon) [1 + \frac{i}{2} \Gamma_{\tilde{\sigma}} \langle \langle d_{\tilde{\sigma}} | d_{\tilde{\sigma}}^{\dagger} \rangle \rangle_{\varepsilon}^{a}]}{E - \varepsilon - \varepsilon_{1} + i \tilde{\gamma}_{1}^{\tilde{\sigma}}}, \quad (B6)$$

$$\Sigma_{2\bar{\sigma}}^{T}(E) = \int \frac{d\varepsilon}{2\pi} \frac{\sum_{\lambda} \Gamma_{\bar{\sigma}}^{\lambda} f_{\lambda}(\varepsilon) [1 - \frac{i}{2} \Gamma_{\bar{\sigma}} \langle \langle d_{\bar{\sigma}} | d_{\bar{\sigma}}^{\dagger} \rangle \rangle_{\varepsilon}^{r}]}{E + \varepsilon - \varepsilon_{2} + i\tilde{\gamma}_{2}}.$$
 (B7)

In the above, we have introduced $\varepsilon_1 = \tilde{\varepsilon}_{\sigma} - \tilde{\varepsilon}_{\bar{\sigma}}$, and $\varepsilon_2 = \tilde{\varepsilon}_{\sigma} + \tilde{\varepsilon}_{\bar{\sigma}} + U$. The subscripts 1 and 2 refer to the excited 1- and 2-electron states of the dot, respectively. The symbols a/r denote advanced/retarded Green function. The self-consistency requires that input dot occupation $\langle n_{\bar{\sigma}} \rangle$ equals that obtained from $G_{\bar{\sigma}}^r(E)$ in the consecutive iteration step with a given accuracy. As noted earlier, there exists exact relation

$$\langle n_{\sigma} \rangle = \int dE \frac{\sum_{\lambda} \Gamma_{\sigma}^{\lambda} f_{\lambda}(E)}{\sum_{\lambda} \Gamma_{\sigma}^{\lambda}} \left(-\frac{1}{\pi} \right) \operatorname{Im} G_{\sigma}^{r}(E), \qquad (B8)$$

valid for energy independent couplings; $\Gamma_{\sigma}^{\lambda}(E) \equiv \Gamma_{\sigma}^{\lambda}$. If this condition is violated, as it might be the case in graphene [48–50], hybrid systems with one (or both) of the electrodes being a superconductor, e.g., *d*-wave [51] one, other approaches are needed.

The inverse lifetimes $\tilde{\gamma}_{\alpha}$ of the excited states $\alpha = |\sigma\rangle$, $|\uparrow\rangle$, $\downarrow\rangle$ stem from higher order processes [42,46]. They can be calculated up to the desired order via the generalized Fermi rule as

$$\tilde{\gamma}_{\alpha} = 2\pi \sum_{|f\rangle} |\langle T(E_{\alpha}) \rangle|^2 \delta(E_{\alpha} - E_f), \tag{B9}$$

with $T(E) = \hat{V} + \hat{V}g(E)\hat{V} + \cdots$ the scattering matrix, where \hat{V} denotes the part of the Hamiltonian describing the coupling between quantum dot and reservoirs. In the discussed approach, one also replaces ε_d by $\tilde{\varepsilon}_d$, to be calculated self-consistently from

$$\tilde{\varepsilon}_d = \varepsilon_d + \Sigma_{1\sigma}^T (\tilde{\varepsilon}_d) + \Sigma_{2\sigma}^T (\tilde{\varepsilon}_d).$$
(B10)

Finally, the self-energies $\Sigma_{\sigma}^{(1,2)}$ are equal to $\Sigma_{0\sigma}$ for $i\tilde{\gamma}_{1,2}^{\alpha} = i0^+$; however, for arbitrary values of $i\tilde{\gamma}_{1,2}^{\alpha}$ they have to be calculated directly from

$$\Sigma_{\sigma}^{(1,2)}(E) = \sum_{\lambda} \Gamma_{\bar{\sigma}}^{\lambda} \int \frac{d\varepsilon}{2\pi} \frac{1}{E \mp \varepsilon - \varepsilon_{1,2} + i\tilde{\gamma}_{1,2}^{\bar{\sigma}}}.$$
 (B11)

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The on-dot density of states is obtained from the retarded Green function. It is defined by Eq. (A6) and shown in Fig. 9 as a function of energy for a number of values of the asymmetry factor α (cf. its definition in Sec. III affecting couplings between the dot and leads). One observes three Abrikosov-Suhl resonances pinned to Fermi levels of the left μ_L , right μ_R and up and down $\mu_U = \mu_D = \mu = 0$ electrodes. Terminal R with $\mu_R = -1.0$ is decoupled from the dot for $\alpha = 0$ so the corresponding density of states shows two Kondo resonances: one at $\mu_U = 0$ and other at $\mu_L = +1.0$. With increasing α the third Kondo resonance appears around energy $E = -1 = \mu_R$. It has to be noted that the Kondo peak at zero energy and two such structures at $V_{L/R}$ make the dependence of currents on gate voltage more complicated in comparison to the two terminal case as the central resonance is always present and only its weight changes with parameters.

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