

Time-dependent quantum transport: A practical scheme using density functional theory

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We present a computationally tractable scheme of time-dependent transport phenomena within open-boundary time-dependent density functional theory. Within this approach all the response properties of a system are determined from the time propagation of the set of “occupied” Kohn-Sham orbitals under the influence of the external bias. This central idea is combined with an open-boundary description of the geometry of the system that is divided into three regions: left/right leads and the device region (“real simulation region”). We have derived a general scheme to extract the set of initial states in the device region that will be propagated in time with proper transparent boundary-condition at the device/lead interface. This is possible due to a new modified Crank-Nicholson algorithm that allows an efficient time-propagation of open quantum systems. We illustrate the method in one-dimensional model systems as a first step towards a full first-principles implementation. In particular we show how a stationary current develops in the system independent of the transient-current history upon application of the bias. The present work is ideally suited to study ac transport and photon-induced charge-injection. Although the implementation has been done assuming clamped ions, we discuss how it can be extended to include dissipation due to electron-phonon coupling through the combined simulation of the electron-ion dynamics as well as electron-electron correlations.

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I. INTRODUCTION

During the last decades, the size of electronic circuits has continuously been reduced. Today, systems like quantum wires and quantum dots are routinely produced on the nanometer scale. Recently, the seemingly ultimate limit of minituarization has been achieved by several experimental groups who were able to place single molecules between two macroscopic electrodes.^{1,2} To describe transport properties on such a small scale, a quantum theory of transport is required.^{3,4}

A cornerstone of such a theory is the Landauer-Büttiker formalism,^{5,6} which provides a method to compute the steady-state current of non-interacting electrons for meso- or nanoscopic systems connecting two (or more) macroscopic electrodes.

Alternatively, the technique of nonequilibrium Green functions (NEG)^{7,8} has been used to tackle quantum transport. Studies using the NEG approach typically use tight-binding-like model Hamiltonians to describe the combined system electrodes plus “device.” A well-known scheme is the one introduced by Caroli *et al.*^{9,10} In the remote past the left and right electrodes are disconnected and in equilibrium at two different chemical potentials; the conducting part of the Hamiltonian is switched on adiabatically and eventually a steady state develops. Within this contacting approach also time-dependent transport phenomena have been investigated.¹¹ Caroli *et al.* discussed non-interacting systems only. Their approach has later been extended to account for short-range electron-electron interaction and for interaction with vibrations in the device region.¹² An excellent overview of the field can be found in the book by Haug and Jauho¹³ and in Ref. 4. Despite its appeal, the above

scheme has limitations since the time-dependent perturbation is the tunneling Hamiltonian, connecting the electrodes to the device, rather than the external electric field.

Cini proposed another scheme¹⁴ also based on NEG. Here, the system electrodes plus “device” is connected and in equilibrium in the remote past. The time-dependent perturbation is the external scalar potential. It has been shown¹⁵ that for non-interacting systems the contacting approach and the Cini approach yield the same current in the long-time limit and that in the dc case the steady-state current does not depend on the history of the applied potential. Moreover, the Cini scheme is well suited for a density functional extension since the electrons are driven out of equilibrium by a local potential rather than by a nonlocal one (see below).

With recent experimental progress to place single molecules as devices between macroscopic electrodes there also has been considerable activity to describe transport through these systems on an *ab initio* level. Most approaches are based on a self-consistency procedure first proposed by Lang.¹⁶ In this steady-state approach based on density functional theory (DFT), exchange and correlation is approximated by the static Kohn-Sham (KS) potential and the charge density is obtained self-consistently in the presence of the steady current. However, the original justification involved subtle points such as different Fermi levels deep inside the left and right electrodes and the implicit reference of nonlocal perturbations such as tunneling Hamiltonians within a DFT framework. (For a detailed discussion we refer to Ref. 15.) The steady-state DFT approach has been further developed^{17–20} and the results have been most useful for understanding the qualitative behavior of measured current-voltage characteristics. Quantitatively, however, the theoretical *I-V* curves often differ from the experimental ones by

several orders of magnitude.²¹ Several explanations are possible for such a mismatch: models are not sufficiently refined, parasitic effects in measurements have been underestimated, and the characteristics of the molecule-contact interface are not well understood and difficult to address given their atomistic complexity. Adding to the theoretical reason for this discrepancy is the fact that the transmission functions computed from static DFT have resonances at the non-interacting Kohn-Sham excitation energies, which in general do not coincide with the true excitation energies. Furthermore, different exchange-correlation functionals lead to DFT currents that vary by more than an order of magnitude.^{21,22}

Excitation energies of interacting systems are accessible via time-dependent (TD) DFT.^{23,24} In this theory, the time-dependent density of an interacting system moving in an external, time-dependent local potential can be calculated via a fictitious system of non-interacting electrons moving in a local, effective time-dependent potential. Therefore this theory is in principle well suited for the treatment of non-equilibrium transport problems.²⁵ A basic issue is that most exchange-correlation functionals have been derived under equilibrium conditions and their application to nonequilibrium problems should be analyzed in more detail. However, this is beyond the scope of the present work.

Before a TDDFT calculation of transport can be tackled, a number of technical problems have to be addressed. In particular, one needs a practical scheme for the propagation of the time-dependent Schrödinger equation for an infinitely large system. Of course, since one can in practice only deal with finite systems, this can only be achieved by applying the correct boundary conditions. The problem of so-called “transparent boundary conditions” for the time-dependent Schrödinger equation has been attacked by many authors. For a recent overview, the reader is referred to Ref. 26.

In this paper we present a propagation scheme that is particularly designed to be used for the calculation of time-dependent transport problems. In Sec. II, we combine the Cini scheme with TDDFT and we develop a general formalism based on the propagation of Kohn-Sham orbitals in open systems. In Sec. III we will address the question of how to obtain the correct initial states for the propagation. An algorithm for the time-evolution of open systems is proposed in Sec. IV. It is based on a modified version of the Crank-Nicholson algorithm. Section V describes some details of our numerical implementation and Sec. VI gives results for several one-dimensional model systems. We draw our conclusions in Sec. VII.

II. GENERAL FORMULATION

We consider an electrode-junction-electrode system that is initially in equilibrium ($t < 0$). The system is contacted and no current flows through the junction, the charge density of the electrodes being perfectly balanced. Therefore, the system initially is in its ground state which, due to the Hohenberg-Kohn theorem,²⁷ is a functional of the density. This density can then be computed in the usual way by $n(\mathbf{r}, 0) = \sum_{occ} |\psi_s(\mathbf{r}, 0)|^2$ where the sum is over the occupied

Kohn-Sham orbitals $\psi_s(\mathbf{r}, 0)$, i.e., the eigenfunctions of the Kohn-Sham Hamiltonian $\mathbf{H}(0)$ with eigenenergy below the Fermi energy. Here and in the following we use boldface notation to denote operators in one-electron Hilbert space.

To observe a current we drive the system out of equilibrium by exposing the electrons to an external electric potential (bias). Without loss of generality we will assume that this external bias vanishes for times $t \leq 0$. According to the Runge-Gross theorem,²³ the time-dependent density can be calculated by evolving the KS orbitals according to the KS equation of TDDFT $i\dot{\psi}_s(t) = \mathbf{H}(t)\psi_s(t)$ where $\mathbf{H}(t)$ is the time-dependent KS Hamiltonian.²⁸ Thus, $n(\mathbf{r}, t) = \sum_{occ} |\psi_s(\mathbf{r}, t)|^2$ and the continuity equation is $(\partial/\partial t)n(\mathbf{r}, t) = -\nabla \cdot \mathbf{j}_{KS}(\mathbf{r}, t)$, where $\mathbf{j}_{KS}(\mathbf{r}, t) = -\sum_{occ} \text{Im}[\psi_s^*(\mathbf{r}, t) \nabla \psi_s(\mathbf{r}, t)]$ is the KS current density. Due to the Runge-Gross theorem and the continuity equation one can deduce that the longitudinal part of the KS current density equals the longitudinal part of the true current density. This need not be true for the transverse part of the current density. However, the transverse part of the current density does not contribute to the total current,²⁹ which can then be calculated by a surface integral

$$I_S(t) = e \sum_{s=1}^N \int_S d\sigma \hat{\mathbf{n}} \cdot \text{Im}[\psi_s^*(\mathbf{r}, t) \nabla \psi_s(\mathbf{r}, t)], \quad (1)$$

where $\hat{\mathbf{n}}$ is the unit vector perpendicular to the surface element $d\sigma$ and the surface S is perpendicular to the longitudinal geometry of our system. In order to propagate the KS orbitals we need to solve the Schrödinger equation for a macroscopic and nonperiodic system. This goal is hopeless unless we know the dynamics of the remote parts of the system. We restrict ourselves to metallic electrodes. Then, the external potential and the disturbance introduced by the device region are screened and the density deep inside the electrodes is equal to the bulk density. As the system size increases, the remote parts are less disturbed by the junction and the density inside the electrodes approaches the equilibrium bulk-density. Thus, the macroscopic size of the electrodes leads to an enormous simplification since the initial-state self-consistency is not disturbed far away from the constriction.

It is convenient to partition the system into three main regions: a central region C consisting of the junction and a few atomic layers of the left and right electrodes and two regions L, R that describe the left and right bulk electrodes. Only the central device region C will be treated explicitly. Our scheme accounts for the full dynamical screening in the central region. It can be further refined by taking into account screening effects also deeper in the electrodes at the level of linear response, with a limited increase in numerical efforts. (These effects might be of importance in the initial transient phase where long-range plasma oscillations in the electrodes may occur.) According to the above partitioning, the original KS Hamiltonian can be written as a 3×3 block matrix, and the Schrödinger equation reads

$$i\frac{\partial}{\partial t}\begin{bmatrix}\psi_L \\ \psi_C \\ \psi_R\end{bmatrix}=\begin{bmatrix}\mathbf{H}_{LL} & \mathbf{H}_{LC} & 0 \\ \mathbf{H}_{CL} & \mathbf{H}_{CC} & \mathbf{H}_{CR} \\ 0 & \mathbf{H}_{RC} & \mathbf{H}_{RR}\end{bmatrix}\begin{bmatrix}\psi_L \\ \psi_C \\ \psi_R\end{bmatrix}, \quad (2)$$

where ψ_α is the projected wave function onto the region $\alpha = L, R, C$. We can solve the differential equation for ψ_L and ψ_R by introducing the retarded Green function \mathbf{g}_α for electrode α , which satisfies $[id/dt - \mathbf{H}_{\alpha\alpha}(t)]\mathbf{g}_\alpha(t, t') = \delta(t - t')$, $\alpha = L, R$, with boundary conditions $\mathbf{g}_\alpha(t^+, t) = -i$ and $\mathbf{g}_\alpha(t, t^+) = 0$. Then, we have for $\alpha = L, R$

$$\psi_\alpha(t) = i\mathbf{g}_\alpha(t, 0)\psi_\alpha(0) + \int_0^t dt' \mathbf{g}_\alpha(t, t')\mathbf{H}_{\alpha C}\psi_C(t'). \quad (3)$$

Using Eq. (3), the equation for ψ_C can be written as

$$i\frac{\partial}{\partial t}\psi_C(t) = \mathbf{H}_{CC}(t)\psi_C(t) + \int_0^t dt' \mathbf{\Sigma}(t, t')\psi_C(t') + i \sum_{\alpha=L,R} \mathbf{H}_{C\alpha}\mathbf{g}_\alpha(t, 0)\psi_\alpha(0), \quad (4)$$

where $\mathbf{\Sigma} = \sum_{\alpha=L,R} \mathbf{H}_{C\alpha}\mathbf{g}_\alpha\mathbf{H}_{\alpha C}$ is the self-energy that accounts for the hopping in and out of region C . Thus, for any given KS orbital we can evolve its projection onto the central region by solving Eq. (4) in region C . Equation (4) has also been derived elsewhere (for static Hamiltonians).³⁰ To summarize, all the complexity of the infinite electrode-junction-electrode system has been reduced to the solution of an open quantum-mechanical system (the central region C) with proper time-dependent boundary conditions.

III. COMPUTATION OF EXTENDED EIGENSTATES

Equation (4) of the previous section is the central equation of our approach to time-dependent transport. It is a reformulation of the original time-dependent Schrödinger equation (2) of the full system in terms of an equation for the central (device) region only. The coupling to the leads is taken into account by the lead Green functions \mathbf{g}_α , $\alpha = L, R$. Equation (4) has the structure of a time-dependent Schrödinger equation with two extra terms: the first term involves the self energy $\mathbf{\Sigma}$, which we will also denote as the memory integral since it involves the wave function in the central region at previous times during the propagation.^{31,32} The second term describes the injection of particles induced by a nonvanishing projection of the initial wave function onto the leads.

Equation (4) is first order in time, therefore we need to specify an initial state that is to be propagated. We want to study the time evolution of systems perturbed out of their equilibrium ground state. Of course, the ground state of our non-interacting system is the Slater determinant of the occupied eigenstates of the full, extended Hamiltonian in equilibrium, $\mathbf{H}(t < 0) = \mathbf{H}^s$. The practical question then arises how one can obtain these eigenstates without having to deal explicitly with the extended Hamiltonian. Note that, unlike in a bulk solid, the translational symmetry is broken in the present device situation.

In the present section we propose a solution of this problem that is based on the partitioning approach used in many

steady-state transport calculations (see, e.g., Ref. 33). The retarded Green function of the static Hamiltonian in the energy domain is determined by

$$[(E + i\eta)\mathbf{1} - \mathbf{H}^s]\mathcal{G}(E) = \mathbf{1}. \quad (5)$$

The Green function $\mathcal{G}(E)$ of the full system can be written in the same block structure as the Hamiltonian

$$\mathcal{G}(E) = \begin{bmatrix} \mathcal{G}_{LL}(E) & \mathcal{G}_{LC}(E) & \mathcal{G}_{LR}(E) \\ \mathcal{G}_{CL}(E) & \mathcal{G}_{CC}(E) & \mathcal{G}_{CR}(E) \\ \mathcal{G}_{RL}(E) & \mathcal{G}_{RC}(E) & \mathcal{G}_{RR}(E) \end{bmatrix}. \quad (6)$$

Equation (5) can be solved for the block of the Green function referring only to the central region

$$\mathcal{G}_{CC}(E) = \frac{1}{(E + i\eta)\mathbf{1}_C - \mathbf{H}_{CC}^s - \sum_{\alpha=L,R} \mathbf{H}_{C\alpha}^s \mathbf{g}_\alpha(E) \mathbf{H}_{\alpha C}^s}, \quad (7)$$

with the retarded Green function of lead α

$$\mathbf{g}_\alpha(E) = \frac{1}{(E + i\eta)\mathbf{1}_\alpha - \mathbf{H}_{\alpha\alpha}^s} \quad (8)$$

and the unit matrix $\mathbf{1}_\alpha$ in region α . This Green function enters as a central ingredient into the Fisher-Lee relation³⁴ for the calculation of the transmission function. Through the coupling to the leads it provides for level broadening of the isolated central part, but it also contains information on the eigenstates of the extended system.

In order to illustrate the central idea of our method to extract the extended eigenstates from the Green function we consider \mathbf{H}^s to be the discretized form of a continuous Hamiltonian $\hat{H}^s(\mathbf{r})$. The continuous Green function and the discretized one for uniform lattice spacing Δx are connected by

$$\mathcal{G}(\mathbf{r}_i, \mathbf{r}_j, E) = \mathcal{G}(E)_{ij} / (\Delta x)^N, \quad (9)$$

where $N=1, 2, 3$ is the number of spatial dimensions of the problem. We choose the convention that a single-particle orbital ψ_{Ej} of the Hamiltonian \hat{H}^s is uniquely specified by its eigenenergy E and a label j for the d_E degenerate orbitals of this energy. Using the Lehmann representation and assuming that \hat{H}^s is invariant under time reversal, the imaginary part of \mathcal{G} is

$$-\frac{1}{\pi} \text{Im}[\mathcal{G}(\mathbf{r}, \mathbf{r}', E)] = \sum_{E'} \delta(E - E') \sum_{j=1}^{d_{E'}} \psi_{E'j}(\mathbf{r}) \psi_{E'j}^*(\mathbf{r}'). \quad (10)$$

Multiplying Eq. (10) by $\psi_{E'}(\mathbf{r}')$, integrating over \mathbf{r}' , using the orthogonality of the single particle states, and dividing by the density of states $D(E) = \sum_{E'} \delta(E - E') d_{E'}$, we obtain

$$-\frac{1}{\pi D(E)} \int d^N r \operatorname{Im}[\mathcal{G}(\mathbf{r}, \mathbf{r}', E)] \psi_{El}(\mathbf{r}') = \gamma(E) \psi_{El}(\mathbf{r}) \quad (11)$$

with

$$\gamma(E) = \frac{1}{D(E)} \sum_{E'} \delta(E - E'). \quad (12)$$

Equation (11) has the structure of an eigenvalue equation where the energy eigenstate ψ_{El} is also an eigenstate of the integral operator $-\operatorname{Im}[\mathcal{G}(\mathbf{r}, \mathbf{r}', E)]/[\pi D(E)]$ with eigenvalue $\gamma(E)$. For a given energy E , this integral operator has d_E different degenerate eigenstates.

We note that Eq. (10) is valid for *all* points in space, in particular also for both \mathbf{r} and \mathbf{r}' representing points in the central region. In this case we know the Green function \mathcal{G}_{CC} through Eqs. (7) and (9). Below we show that the eigenfunctions of $\operatorname{Im}(\mathcal{G}_{CC})$ can be expressed as linear combination of the ψ_{El} . Let us consider the matrix formed by the elements

$$b_{ml} = \int_C d^N r \psi_{Em}^*(\mathbf{r}) \psi_{El}(\mathbf{r}), \quad (13)$$

where the integration is over the central region only. This matrix is Hermitian and can be diagonalized, i.e.,

$$\sum_{l=1}^{d_E} b_{ml} a_{jl} = \lambda_j a_{jm} \quad (14)$$

with λ_j real. Next we compute the matrix elements of the Green function with respect to the functions

$$a_{Ej}(\mathbf{r}) = \sum_{l=1}^{d_E} a_{jl}(E) \psi_{El}(\mathbf{r}). \quad (15)$$

After a straightforward manipulation one finds

$$\begin{aligned} & -\frac{1}{\pi} \int_C d^N r \int_C d^N r' a_{El}^*(\mathbf{r}) \operatorname{Im}[\mathcal{G}(\mathbf{r}, \mathbf{r}', E)] a_{Ej}(\mathbf{r}') \\ & = \delta_{jl} \lambda_j^2 \sum_{E'} \delta(E - E'), \end{aligned} \quad (16)$$

which shows explicitly that the functions $a_{Ej}(\mathbf{r})$ diagonalize $\operatorname{Im}(\mathcal{G}_{CC})$ in the central region and that the eigenvalues are positive. Since any linear combination of degenerate eigenstates is again an eigenstate, diagonalizing $\operatorname{Im}[\mathcal{G}_{CC}(E)]$ gives us one set of linearly independent, degenerate eigenstates of energy E .

In our practical implementation described in more detail in Sec. V, we diagonalize

$$-\frac{1}{\pi D_C(E)} \operatorname{Im}[\mathcal{G}_{CC}(E)], \quad (17)$$

where

$$D_C(E) = -\frac{1}{\pi} \operatorname{Tr}\{\operatorname{Im}[\mathcal{G}_{CC}(E)]\} \quad (18)$$

is the total density of states in the central region. If we use N_g grid points to describe the central region, the diagonalization in principle gives N_g eigenvectors but only a few have the physical meaning of extended eigenstates at this energy. It is, however, very easy to identify the physical states by looking at the eigenvalues: only a few eigenvalues (for the simple examples we studied either one or two) are nonvanishing and they always add up to unity. The corresponding states are the physical ones. All the other eigenvalues are zero (or numerically close to zero) and the corresponding states have no physical meaning.

The procedure described above gives the correct extended eigenstates only up to a normalization factor. When diagonalizing Eq. (17) with typical library routines one obtains eigenvectors that are normalized to the central region. Physically this might be incorrect. Therefore, the normalization has to be fixed separately. In the example of Sec. V we fixed the norm by matching the wave function for the central region to the known form (and normalization) of the wave function in the macroscopic leads.

It should be emphasized that the procedure described here for the extraction of eigenstates of the extended system from $\mathcal{G}_{CC}(E)$ only works in practice if E is in the continuous part of the spectrum due to the sharp peak of the delta function in the discrete part of the spectrum.

Eigenstates in the discrete part of the spectrum can be found by considering the original Schrödinger equation for the full system:

$$\mathbf{H}^s \psi = E \psi. \quad (19)$$

Using again the block structure of the Hamiltonian, this can be transformed into an effective Schrödinger equation for an *energy-dependent* Hamiltonian for the central region only:

$$\left(\mathbf{H}_{CC}^s - \sum_{\alpha=L,R} \mathbf{H}_{C\alpha} \mathbf{g}_\alpha(E) \mathbf{H}_{\alpha C} \right) \psi_C = E \psi_C. \quad (20)$$

This equation has solutions only for certain values of E , which are the discrete eigenenergies of the full Hamiltonian \hat{H}^s . Therefore, one can find these states by iteration. We have successfully tested this idea for systems where the analytic solutions are known. However, since the main focus of the present work is transport where the continuum states are the essential physical ingredient, we will not deal with the states in the discrete spectrum for the remainder of this paper. Those states might play a role in the description of charge accumulation in molecular transport, as, e.g., in Coulomb blockade phenomena.

IV. ALGORITHM FOR TIME EVOLUTION

In order to calculate the longitudinal current in an electrode-junction-electrode system we need to propagate the Kohn-Sham orbitals. The main difficulty stems from the macroscopic size of the electrodes whose remote parts, ultimately, are taken infinitely far away from the central, explicitly treated, scattering region C .

The problem can be solved by imposing transparent boundary conditions²⁶ at the electrode-junction interfaces. Efficient algorithms have been proposed for wave packets initially *localized* in the scattering region and for Hamiltonians constant in time. In this section we propose an algorithm well suited for delocalized initial states, as well as for localized ones, evolving with a time-dependent Hamiltonian.

Let $\mathbf{H}(t)$ be the time-dependent KS Hamiltonian. We partition $\mathbf{H}(t)$ as in Sec. II. The explicitly treated region C includes the first few atomic layers of the left and right electrodes. The boundaries of this region are chosen in such a way that the density outside C is accurately described by an equilibrium bulk density. It is convenient to write $\mathbf{H}_{\alpha\alpha}(t)$, with $\alpha=L,R$, as the sum of a term $\mathbf{H}_{\alpha\alpha}^s$, which is constant in time, and another term $\mathbf{U}_\alpha(t)$, which is explicitly time-dependent, $\mathbf{H}_{\alpha\alpha}(t)=\mathbf{H}_{\alpha\alpha}^s+\mathbf{U}_\alpha(t)$. In configuration space $\mathbf{U}_\alpha(t)$ is diagonal at any time t since the KS potential is local in space. Furthermore, the diagonal elements $U_\alpha(\mathbf{r},t)$ are spatially constant for metallic electrodes. Thus, $\mathbf{U}_\alpha(t)=U_\alpha(t)\mathbf{1}_\alpha$ and $U_L(t)-U_R(t)$ is the total potential drop across the central region. We write $\mathbf{H}(t)=\tilde{\mathbf{H}}(t)+\mathbf{U}(t)$ with

$$\tilde{\mathbf{H}}(t)=\begin{bmatrix} \mathbf{H}_{LL}^s & \mathbf{H}_{LC} & 0 \\ \mathbf{H}_{CL} & \mathbf{H}_{CC}(t) & \mathbf{H}_{CR} \\ 0 & \mathbf{H}_{RC} & \mathbf{H}_{RR}^s \end{bmatrix}, \quad (21)$$

and

$$\mathbf{U}(t)=\begin{bmatrix} U_L(t)\mathbf{1}_L & 0 & 0 \\ 0 & 0 & 0 \\ 0 & 0 & U_R(t)\mathbf{1}_R \end{bmatrix}. \quad (22)$$

In this way, the only term in $\tilde{\mathbf{H}}(t)$ that depends on t is $\mathbf{H}_{CC}(t)$. For any given initial state $\psi(0)=\psi^{(0)}$ we calculate $\psi(t_m=m\Delta t)=\psi^{(m)}$ by using a generalized form of the Cayley method

$$(1+i\delta\tilde{\mathbf{H}}^{(m)})\frac{1+i\frac{\delta}{2}\mathbf{U}^{(m)}}{1-i\frac{\delta}{2}\mathbf{U}^{(m)}}\psi^{(m+1)}=(1-i\delta\tilde{\mathbf{H}}^{(m)})\frac{1-i\frac{\delta}{2}\mathbf{U}^{(m)}}{1+i\frac{\delta}{2}\mathbf{U}^{(m)}}\psi^{(m)}, \quad (23)$$

with $\tilde{\mathbf{H}}^{(m)}=\frac{1}{2}[\tilde{\mathbf{H}}(t_{m+1})+\tilde{\mathbf{H}}(t_m)]$, $\mathbf{U}^{(m)}=\frac{1}{2}[\mathbf{U}(t_{m+1})+\mathbf{U}(t_m)]$, and $\delta=\Delta t/2$. It should be noted that our propagator is norm conserving (unitary) and accurate to second order in δ , as is the Cayley propagator.³⁵ Denoting by ψ_α the projected wave function onto the region $\alpha=R,L,C$, we find from Eq. (23)

$$\psi_C^{(m+1)}=\frac{1-i\delta\mathbf{H}_{\text{eff}}^{(m)}}{1+i\delta\mathbf{H}_{\text{eff}}^{(m)}}\psi_C^{(m)}+S^{(m)}-M^{(m)}. \quad (24)$$

Here, $\mathbf{H}_{\text{eff}}^{(m)}$ is the effective Hamiltonian of the central region:

$$\mathbf{H}_{\text{eff}}^{(m)}=\mathbf{H}_{CC}^{(m)}-i\delta\mathbf{H}_{CL}(1+i\delta\mathbf{H}_{LL}^s)^{-1}\mathbf{H}_{LC}-i\delta\mathbf{H}_{CR}(1+i\delta\mathbf{H}_{RR}^s)^{-1}\mathbf{H}_{RC}$$

with $\mathbf{H}_{CC}^{(m)}=\frac{1}{2}[\mathbf{H}_{CC}(t_{m+1})+\mathbf{H}_{CC}(t_m)]$. The source term $S^{(m)}$ describes the injection of density into the region C , while the

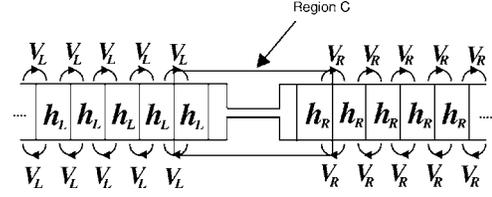


FIG. 1. The schematic sketch of an electrode-junction-electrode system with semiperiodic electrodes.

memory term $M^{(m)}$ is responsible for the hopping in and out of the region C . In terms of the propagator for the uncontacted and undisturbed α electrode

$$\mathbf{g}_\alpha=\frac{1-i\delta\mathbf{H}_{\alpha\alpha}^s}{1+i\delta\mathbf{H}_{\alpha\alpha}^s}, \quad (25)$$

the source term can be written as

$$S^{(m)}=-\frac{2i\delta}{1+i\delta\mathbf{H}_{\text{eff}}^{(m)}}\sum_{\alpha=L,R}\frac{\Lambda_\alpha^{(m,0)}}{u_\alpha^{(m)}}\mathbf{H}_{C\alpha}\frac{(\mathbf{g}_\alpha)^m}{1+i\delta\mathbf{H}_{\alpha\alpha}^s}\psi_\alpha^{(0)}, \quad (26)$$

with

$$u_\alpha^{(m)}=\frac{1-i\frac{\delta}{2}\mathbf{U}_\alpha^{(m)}}{1+i\frac{\delta}{2}\mathbf{U}_\alpha^{(m)}} \text{ and } \Lambda_\alpha^{(m,k)}=\prod_{j=k}^m (u_\alpha^{(j)})^2. \quad (27)$$

For a wave packet initially localized in C the projection onto the left and right electrode $\psi_\alpha^{(0)}$ vanishes and $S^{(m)}=0$ for any m , as it should be. The memory term is more complicated and reads

$$M^{(m)}=-\frac{\delta^2}{1+i\delta\mathbf{H}_{\text{eff}}^{(m)}}\sum_{\alpha=L,R}\sum_{k=0}^{m-1}\frac{\Lambda_\alpha^{(m,k)}}{u_\alpha^{(m)}u_\alpha^{(k)}}(\mathbf{Q}_\alpha^{(m-k)}+\mathbf{Q}_\alpha^{(m-k-1)})\times(\psi_C^{(k+1)}+\psi_C^{(k)}), \quad (28)$$

where

$$\mathbf{Q}_\alpha^{(m)}=\mathbf{H}_{C\alpha}\frac{[\mathbf{g}_\alpha]^m}{1+i\delta\mathbf{H}_{\alpha\alpha}^s}\mathbf{H}_{\alpha C}. \quad (29)$$

The quantities $\mathbf{Q}_\alpha^{(m)}$ depend on the geometry of the system and are independent of the initial state $\psi^{(0)}$.

The structural similarities of our discretization scheme (24) and Eq. (4) are apparent. The memory term (28) corresponds to the second term on the rhs of Eq. (4), which is nonlocal in time, while the source term (26) corresponds to the last term in Eq. (4). However, our scheme does not follow from a direct discretization of Eq. (4) but rather from a unitary discretization (23) of the time-dependent Schrödinger equation for the extended system.

Below we propose a recursive scheme to calculate the $\mathbf{Q}_\alpha^{(m)}$'s for those system geometries having semiperiodic electrodes along the longitudinal direction (see Fig. 1). In this case $\mathbf{H}_{\alpha\alpha}^s$ has a tridiagonal block form

$$\mathbf{H}_{\alpha\alpha}^s = \begin{bmatrix} \mathbf{h}_\alpha & \mathbf{V}_\alpha & 0 & \dots \\ \mathbf{V}_\alpha & \mathbf{h}_\alpha & \mathbf{V}_\alpha & \dots \\ 0 & \mathbf{V}_\alpha & \mathbf{h}_\alpha & \dots \\ \dots & \dots & \dots & \dots \end{bmatrix}, \quad (30)$$

where \mathbf{h}_α describes a convenient cell and \mathbf{V}_α is the hopping Hamiltonian between two nearest neighbor cells. Without loss of generality we assume that both \mathbf{h}_α and \mathbf{V}_α are square matrices of dimension $N_\alpha \times N_\alpha$. Taking into account that the central region contains the first few cells of the left and right electrodes, the matrix $\mathbf{Q}_\alpha^{(m)}$ has the following structure:

$$\mathbf{Q}_L^{(m)} = \begin{bmatrix} \mathbf{q}_L^{(m)} & 0 & 0 \\ 0 & 0 & 0 \\ 0 & 0 & 0 \end{bmatrix}, \quad \mathbf{Q}_R^{(m)} = \begin{bmatrix} 0 & 0 & 0 \\ 0 & 0 & 0 \\ 0 & 0 & \mathbf{q}_R^{(m)} \end{bmatrix}. \quad (31)$$

The $\mathbf{q}_\alpha^{(m)}$'s are square matrices of dimension $N_\alpha \times N_\alpha$ and are given by

$$\mathbf{q}_\alpha^{(m)} = \mathbf{V}_\alpha \left(\frac{[\mathbf{g}_\alpha]_m}{1 + i\delta\mathbf{H}_{\alpha\alpha}} \right)_{1,1} \mathbf{V}_\alpha, \quad (32)$$

where the subscript (1, 1) denotes the first diagonal block of the matrix in the square brackets. We introduce the generating matrix function

$$\mathbf{q}_\alpha(x, y) \equiv \mathbf{V}_\alpha \left(\frac{1}{x + iy\delta\mathbf{H}_{\alpha\alpha}} \right)_{1,1} \mathbf{V}_\alpha, \quad (33)$$

which can also be expressed in terms of continued matrix fractions (see the Appendix)

$$\mathbf{q}_\alpha(x, y) = \mathbf{V}_\alpha \frac{1}{x + iy\delta\mathbf{h}_\alpha + y^2\delta^2\mathbf{q}_\alpha(x, y)} \mathbf{V}_\alpha. \quad (34)$$

The $\mathbf{q}_\alpha^{(m)}$'s can be obtained from

$$\mathbf{q}_\alpha^{(m)} = \frac{1}{m!} \left(-\frac{\partial}{\partial x} + \frac{\partial}{\partial y} \right)^m \mathbf{q}_\alpha(x, y) \Big|_{x=y=1}. \quad (35)$$

From Eqs. (35) and (34) one can build up a recursive scheme. Let us define $\mathbf{p}_\alpha^{-1}(x, y) = x + iy\delta\mathbf{h}_\alpha + y^2\delta^2\mathbf{q}_\alpha(x, y)$ and $\mathbf{p}_\alpha^{(m)} = 1/m!(-\partial/\partial x + \partial/\partial y)^m \mathbf{p}_\alpha(x, y)|_{x=y=1}$. Then, by definition,

$$\mathbf{q}_\alpha^{(m)} = \mathbf{V}_\alpha \mathbf{p}_\alpha^{(m)} \mathbf{V}_\alpha. \quad (36)$$

Using the identity $1/m!(-\partial/\partial x + \partial/\partial y)^m \mathbf{p}_\alpha(x, y) \mathbf{p}_\alpha^{-1}(x, y) = 0$, one finds

$$\begin{aligned} (1 + i\delta\mathbf{h}_\alpha) \mathbf{p}_\alpha^{(m)} &= (1 - i\delta\mathbf{h}_\alpha) \mathbf{p}_\alpha^{(m-1)} \\ &\quad - \delta^2 \sum_{k=0}^m (\mathbf{q}_\alpha^{(k)} + 2\mathbf{q}_\alpha^{(k-1)} + \mathbf{q}_\alpha^{(k-2)}) \mathbf{p}_\alpha^{(m-k)} \end{aligned} \quad (37)$$

with $\mathbf{p}_\alpha^{(m)} = \mathbf{q}_\alpha^{(m)} = 0$ for $m < 0$. Once $\mathbf{q}_\alpha^{(0)}$ has been obtained by solving Eq. (34) with $x=y=1$, we can calculate $\mathbf{p}_\alpha^{(0)} = (1 + i\delta\mathbf{h}_\alpha + \delta^2\mathbf{q}_\alpha^{(0)})^{-1}$. Afterwards, we can use Eq. (37) with $\mathbf{q}_\alpha^{(0)} = \mathbf{V}_\alpha \mathbf{p}_\alpha^{(1)} \mathbf{V}_\alpha$ to calculate $\mathbf{p}_\alpha^{(1)}$ and hence $\mathbf{q}_\alpha^{(1)}$ and so on and so forth.

This concludes the description of our algorithm for the propagation of the time-dependent Schrödinger equation for

extended systems. It is worth mentioning an additional complication here that arises for the propagation of a time-dependent Kohn-Sham equation. This complication stems from the fact that in order to compute $\psi_C^{(m+1)}$ at time step $m+1$ one needs to know the time-dependent KS potential at the same time step which, via the Hartree and exchange-correlation potentials, depends on the yet unknown orbitals $\psi_C^{(m+1)}$. Of course, the solution is to use a predictor-corrector approach: in the first step one approximates $\mathbf{H}_{CC}^{(m)}$ by $\mathbf{H}_{CC}(t_m)$, computes new orbitals $\tilde{\psi}_C^{(m+1)}$, and from those obtains an improved approximation for $\mathbf{H}_{CC}^{(m)}$.

V. IMPLEMENTATION DETAILS

All the methodological discussion above is general and can be applied to general device configurations as long as they can be mapped into a longitudinal-like geometry as described in Fig. 1. In order to demonstrate the feasibility of the scheme described in the previous sections we have implemented it for one-dimensional model systems. The extension to real molecular-device configurations is presently under development.³⁶ We consider systems described by the Hamiltonian

$$\hat{H} = -\frac{1}{2} \frac{d^2}{dx^2} + V(x). \quad (38)$$

We have used a simple three-point discretization for the second derivative

$$\frac{d^2}{dx^2} \psi(x)|_{x=x_i} \approx \frac{1}{(\Delta x)^2} [\psi(x_{i+1}) - 2\psi(x_i) + \psi(x_{i-1})] \quad (39)$$

with equidistant grid points x_i , $i=1, \dots, N_g$ and spacing Δx . Within this approximation matrices of the form $\mathbf{H}_{C\alpha} \mathbf{M} \mathbf{H}_{\alpha C}$, which are $N_g \times N_g$ matrices and appear, e.g., in Eq. (7) or (29), have only one nonvanishing matrix element. For $\alpha=L$ this is the (1,1) element; for $\alpha=R$ it is the (N_g, N_g) element.

In order to proceed we have to specify the nature of the leads and therefore the lead Green function. Here we choose the simplest case of semi-infinite, uniform leads at constant potential $U_{\alpha 0}$. In this case, the Green function (8) in the energy domain can be given in closed form:

$$\begin{aligned} [\mathbf{g}_\alpha(E)]_{kl} &= -\frac{i\Delta x}{\sqrt{2\tilde{E}_\alpha}} \exp(i\sqrt{2\tilde{E}_\alpha}|x_k - x_l|) \\ &\quad + \frac{i\Delta x}{\sqrt{2\tilde{E}_\alpha}} \exp(i\sqrt{2\tilde{E}_\alpha}(|x_k - x_{\alpha 0}| + |x_l - x_{\alpha 0}|)) \end{aligned} \quad (40)$$

with $\tilde{E}_\alpha = E - U_{\alpha 0}$. The abscissa $x_{\alpha 0}$ is the position of the interface between the lead and the device region and $x_k = x_{\alpha 0} \pm k\Delta x$, where the plus sign applies for $\alpha=R$ and the minus sign for $\alpha=L$.

The results of the procedure for calculating extended eigenstates as described in Sec. III are illustrated in Fig. 2 for a square potential barrier with zero potential in both leads. In the upper panel we have the square modulus of eigenstates at

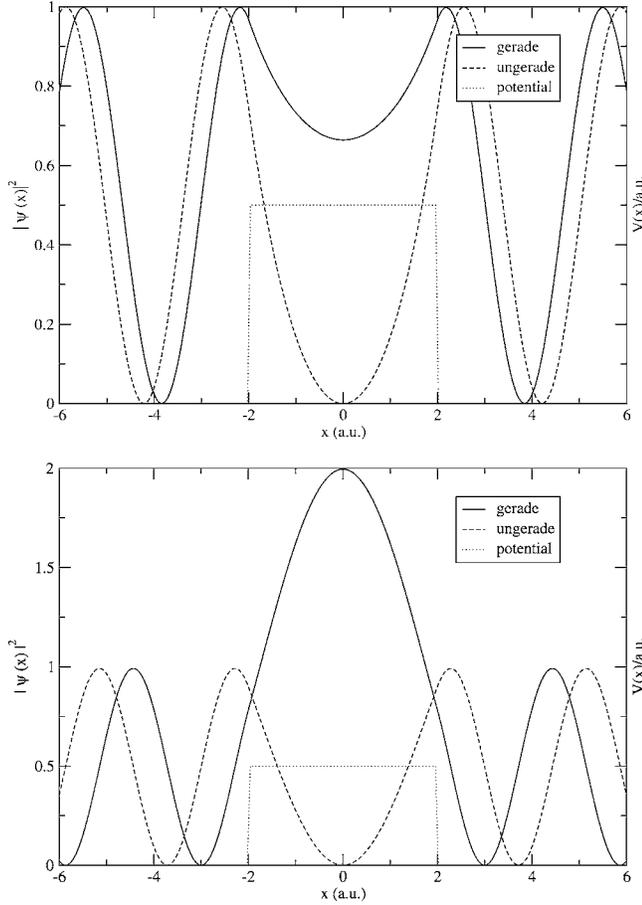


FIG. 2. The continuum states of the square potential barrier at different energies with leads at zero potential. Upper panel: eigenstates for $\varepsilon=0.45$ a.u., just below the barrier height of 0.5 a.u.. Lower panel: eigenstates at $\varepsilon=0.6$ a.u..

an energy below the barrier height while in the lower panel eigenstates with energy higher than the barrier are shown. The states result from diagonalization of Eq. (17). In order to obtain the normalization constant we compute the logarithmic derivative at the boundary of the central region numerically and match it to the analytic form in the lead to obtain the phase shift δ_α :

$$\frac{1}{2} \frac{d^2}{dx^2} \ln[|\psi(x)|^2] \Big|_{x=x_{\alpha 0}} = q \cot(\delta_\alpha) \quad (41)$$

where $q = \sqrt{2\tilde{E}_\alpha}$. Knowing the phase shift, we can rescale the wave function such that it matches with the analytic form $\sin[q(x-x_{\alpha 0}) + \delta_\alpha]$ at the interface. Of course, this form of the extended states only applies for $\tilde{E}_\alpha > 0$, but as long as E is in the continuous part of the spectrum, it is correct at least for one of the leads. This is sufficient to determine the normalization. The states obtained numerically with this procedure coincide with the known analytical results.

We then implemented the propagation scheme presented in the previous section. Within our three-point approximation, \mathbf{h}_α , \mathbf{V}_α , and \mathbf{q}_α are 1×1 matrices. The equation for $q_\alpha^{(0)}$

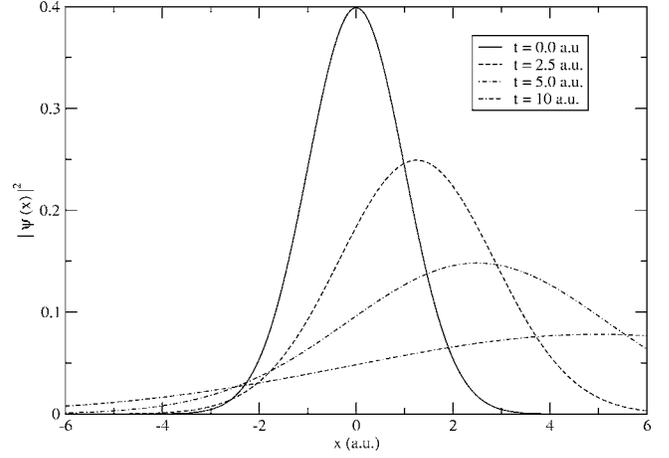


FIG. 3. The time evolution of a Gaussian wave packet with initial width 1.0 a.u. and initial momentum 0.5 a.u. for various propagation times. The transparent boundary conditions allow the wave packet to pass the propagation region without spurious reflections at the boundaries.

[see Eqs. (34) and (35)] becomes a simple quadratic equation that can be solved explicitly:

$$q_\alpha^{(0)} = \frac{-(1 + i\delta h_\alpha) + \sqrt{(1 + i\delta h_\alpha)^2 + 4(\delta V_\alpha)^2}}{2\delta^2}. \quad (42)$$

Although the quadratic equation has two solutions, the above choice for $q_\alpha^{(0)}$ is dictated by the fact that the Taylor expansions for small δ of Eqs. (42) and (34) have to coincide. Using this result we then solved the iterative scheme to obtain the $q_\alpha^{(m)}$ for $m \geq 1$.

As a first check on the propagation method we propagated a Gaussian wave packet which, at initial time $t=0$, is completely localized in the central device region. (The source terms $S^{(m)}$ then vanish identically.) As can be seen in Fig. 3, the wave packet correctly propagates through the boundaries without any spurious reflections.

For the propagation of the extended initial states (the eigenstates of the unperturbed system) we also need to implement the source terms $S^{(m)}$. In the following we assume that the left and right leads are at the same potential initially so that the analytic form of the initial states is in both leads given by $\sin[q(x-x_{\alpha 0}) + \delta_\alpha] = \cos(\delta_\alpha)\sin[q(x-x_{\alpha 0})] + \sin(\delta_\alpha)\cos[q(x-x_{\alpha 0})]$. The propagation of the term proportional to $\sin[q(x-x_{\alpha 0})]$ is trivial since this is an eigenstate of the lead Hamiltonian with energy $\varepsilon_q = q^2/2$. Therefore, if—in discretized form— $\psi_{R,1}^{(0)} = [\sin(q\Delta x), \sin(2q\Delta x), \dots]^T$, we obtain

$$\mathbf{H}_{CR} \frac{[\mathbf{g}_R]^m}{1 + i\delta \mathbf{H}_{RR}^s} \psi_{R,1}^{(0)} = V_R \frac{(1 - i\delta \varepsilon_q)^m}{(1 + i\delta \varepsilon_q)^{m+1}} \mathbf{e}_R \quad (43)$$

and similarly for the left lead. Here, \mathbf{H}_{RR}^s is the static part of the right-lead Hamiltonian, \mathbf{g}_R is the corresponding Green function according to definition (25), and $\mathbf{e}_R = (0, \dots, 0, 1)^T$ is a unit vector.

The propagation of the part proportional to $\cos[q(x-x_{\alpha 0})]$ is more complicated since this is not an eigenstate of

the lead Hamiltonian. We define the function $R(x, y)$ from

$$R(x, y)e_R = \mathbf{H}_{CR} \frac{1}{x + iy \delta \mathbf{H}_{RR}^s} \psi_{R,2}^{(0)}, \quad (44)$$

where $\psi_{R,2}^{(0)} = (\cos(q\Delta x), \cos(2q\Delta x), \dots)^T$. Introducing the tridiagonal matrix

$$\mathbf{O} = \begin{pmatrix} 0 & 1 & 0 & 0 & \cdots \\ 1 & 0 & 1 & 0 & \cdots \\ 0 & 1 & 0 & 1 & \cdots \\ \vdots & \vdots & \vdots & \vdots & \ddots \end{pmatrix} \quad (45)$$

and using

$$V_R \mathbf{O} \psi_{R,2}^{(0)} = (\varepsilon_q - h_R) \psi_{R,2}^{(0)} - V_R(1, 0, 0, \dots)^T \quad (46)$$

one arrives at

$$R(x, y) = \frac{1}{x + iy \delta \varepsilon_q} [V_R \cos(q\Delta x) + iy \delta q_R(x, y)], \quad (47)$$

where $q_R(x, y)$ is given by Eq. (33). Defining

$$R^{(m)} = \frac{1}{m!} \left(-\frac{\partial}{\partial x} + \frac{\partial}{\partial y} \right)^m R(x, y)|_{x=y=1} \quad (48)$$

one finds

$$R^{(m)} = \frac{(1 - i \delta \varepsilon_q)^m}{(1 + i \delta \varepsilon_q)^{m+1}} V_R \cos(q\Delta x) + i \delta \sum_{k=0}^m \frac{(1 - i \delta \varepsilon_q)^{m-k}}{(1 + i \delta \varepsilon_q)^{m+1-k}} (q_R^{(k)} + q_R^{(k-1)}) \quad (49)$$

and finally

$$\mathbf{H}_{CR} \frac{[\mathbf{g}_R]^m}{1 + i \delta \mathbf{H}_{RR}^s} \psi_{R,2}^{(0)} = R^{(m)} e_R. \quad (50)$$

One may proceed along the same lines for extracting the left component.

To test our implementation we have propagated eigenstates of the extended system. As expected, these states just pick up an exponential phase factor $\exp(-iEt)$ during the propagation.

VI. EXAMPLES

We are now in a position to apply our algorithm to the calculation of time-dependent currents in one-dimensional model systems. The systems are initially in thermodynamic equilibrium. At time $t=0$, a bias is switched on in the electrodes.

As a first example we considered a system where the electrostatic potential vanishes identically both in the left and right leads as well as in the central region, which is explicitly propagated. Initially, all single particle levels are occupied up to the Fermi energy ε_F . At $t=0$ a constant bias is switched on in the leads and the time evolution of the system is calculated. We chose the bias in the right lead as the negative of the bias in the left lead, $U_R = -U_L$. The current is calculated from Eq. (1):

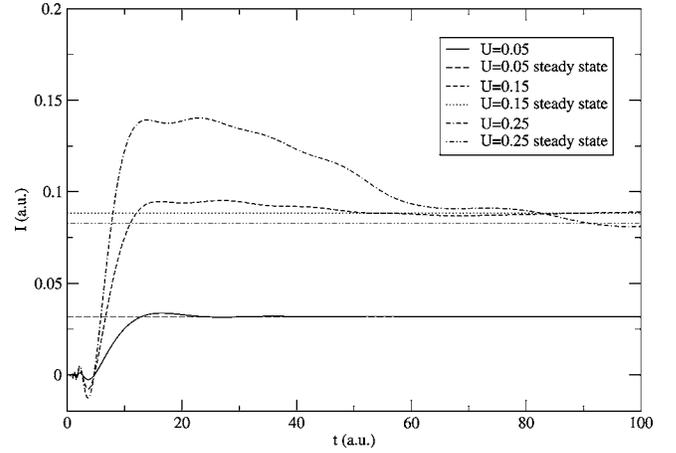


FIG. 4. The time evolution of the current for a system where initially the potential is zero in the leads and the propagation region. At $t=0$, a constant bias with opposite sign in the left and right leads is switched on, $U = U_L = -U_R$ (values in atomic units). The propagation region extends from $x = -6$ to $x = +6$ a.u. The Fermi energy of the initial state is $\varepsilon_F = 0.3$ a.u. The current in the center of the propagation region is shown.

$$I(x, t) = 2 \int_{-k_F}^{k_F} \frac{dk}{2\pi} \text{Im} \left(\psi_k^*(x, t) \frac{d}{dx} \psi_k(x, t) \right) = 2 \int_0^{k_F} \frac{dk}{2\pi} \text{Im} \left(\psi_k^* \frac{d}{dx} \psi_k + \psi_{-k}^* \frac{d}{dx} \psi_{-k} \right), \quad (51)$$

where the prefactor 2 comes from spin and $k_F = \sqrt{2\varepsilon_F}$ is the Fermi wave vector.

The numerical parameters are as follows: the Fermi energy is $\varepsilon_F = 0.3$ a.u., the bias is $U_L = -U_R = 0.05, 0.15, 0.25$ a.u., and the central region extends from $x = -6$ to $x = +6$ a.u. with equidistant grid points with spacing $\Delta x = 0.03$ a.u.. The k integral in Eq. (51) is discretized with 100 k points, which amounts to a propagation of 200 states. The time step for the propagation was $\Delta t = 10^{-2}$ a.u.

In Fig. 4 we have plotted the current densities at $x=0$ as a function of time for different values of the applied bias. As a first feature we notice that a steady state is achieved, in agreement with the results of Ref. 15. The corresponding steady-state current I can be calculated from the Landauer formula. For the present geometry this leads to the steady current

$$I = 8e \int_{\max(U_L, U_R)} \frac{d\omega}{2\pi} [f(\omega - U_L) - f(\omega - U_R)] \times \frac{\sqrt{\omega - U_L} \sqrt{\omega - U_R}}{(\sqrt{\omega - U_L} + \sqrt{\omega - U_R})^2 + U_L U_R \left(\frac{\sin(l\sqrt{2\omega})}{\sqrt{\omega}} \right)^2}, \quad (52)$$

where l is the width of the central region. From Eq. (52) with $l = 12$ a.u. and $U_L = -U_R$, the numerical values for the steady-state currents are 0.0316 a.u. ($U_L = 0.05$ a.u.), 0.0883 a.u. ($U_L = 0.15$ a.u.), and 0.0828 a.u. ($U_L = 0.25$ a.u.). We see that

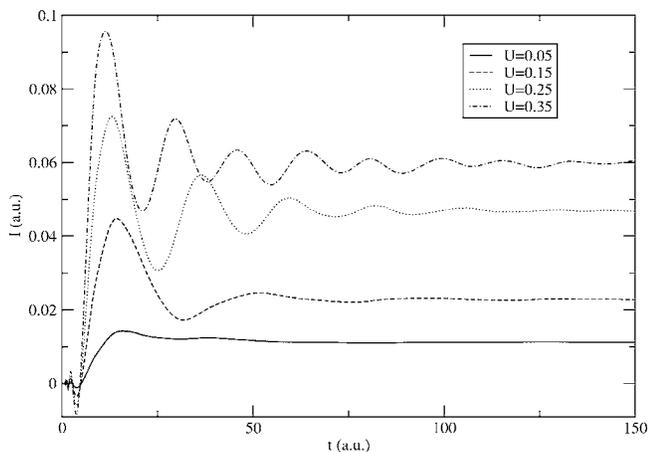


FIG. 5. Time evolution of the current through a double square potential barrier in response to an applied constant bias (given in atomic units) in the left lead. The potential is given by $V(x) = 0.5$ a.u. for $5 \leq |x| \leq 6$ a.u. and zero otherwise; the propagation region extends from $x = -6$ to $x = +6$ a.u. The Fermi energy of the initial state is $\varepsilon_F = 0.3$ a.u. The current in the center of the structure is shown.

our algorithm yields the same answers. Second, we notice that the onset of the current is delayed in relation to the switching time $t=0$. This is easily explained by the fact that the perturbation at $t=0$ happens in the leads only, e.g., for $|x| > 6$ a.u., while we plot the current at $x=0$. In other words, we see the delay time needed for the perturbation to propagate from the leads to the center of our device region. This delay time is given by $t_0 = d/v_F$ where d is the distance from the leads of the point $x=0$ (for which we plot the current) and v_F is the Fermi velocity. We also note that the higher the bias the more the current overshoots its steady-state value for small times after switching on the bias. Finally, it is worth mentioning that increasing the bias does not necessarily lead to a larger steady-state current.

In the second example we studied a double square potential barrier with electrostatic potential $V(x) = 0.5$ a.u. for $5 \text{ a.u.} \leq |x| \leq 6 \text{ a.u.}$ and zero otherwise. This time we switch on a constant bias in the left lead only, i.e., $U_R = 0$. The Fermi energy for the initial state is $\varepsilon_F = 0.3$ a.u.. The central region extends from $x = -6$ to $x = +6$ a.u. with a lattice spacing of $\Delta x = 0.03$ a.u.. Again, we use 100 different k values to compute the current and a time step of $\Delta t = 10^{-2}$ a.u.

In Fig. 5 we plot the current at $x=0$ as a function of time for several values of the applied bias $U = U_L$. Again, a steady state is achieved for all values of U . As discussed in Fig. 4 the transient current can exceed the steady current; the higher the applied voltage, the larger is this excess current and the shorter is the time when it reaches its maximum. Furthermore, the oscillatory evolution towards the steady current solution depends on the bias. For higher bias the frequency of the transient oscillations increases roughly proportional to the applied bias. For small bias the electrons at the bottom of the band are not disturbed and the transient process is exponentially short. On the other hand, for a bias close to the Fermi energy the transient process decays as a power law, due to the band edge singularity. As pointed out in Ref. 15,

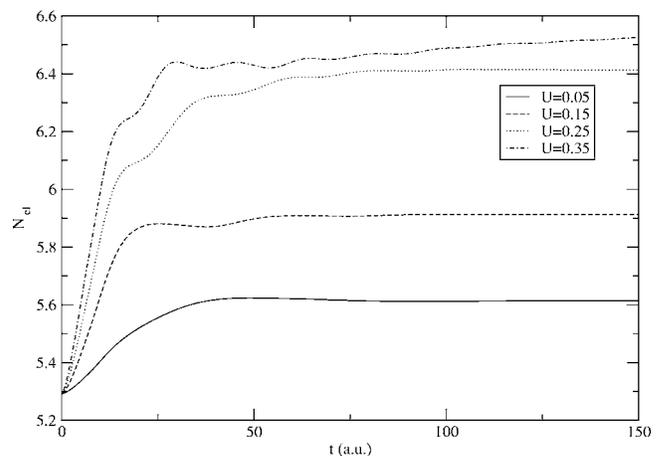


FIG. 6. The time evolution of the total number of electrons in the region $|x| \leq 6$ for the double square potential barrier of Fig. 5.

for non-interacting electrons the steady-state current develops by means of a pure dephasing mechanism. In our examples the transient process occurs in a femtosecond time scale, which is much shorter than the relaxation time due to electron-phonon interactions.

In Fig. 6 we plot the time evolution of the total number of electrons in the device region for the same values of U_L . We see that as a result of the bias a quite substantial amount of charge is added to the device region. This result has important implications when simulating the transport through an interacting system as the effective (dynamical) electronic screening is modified due not only to the external field but also to the accumulation of charge state in the molecular device. This illustrates that linear response might not be an appropriate tool to tackle the dynamical response and that we will need to resort to a full time-propagation approach like the one of the present work. Here we emphasize that all our calculations are done without taking into account the electron-electron interaction. If we had done a similar calculation with the interaction incorporated in a time-dependent Hartree or time-dependent DFT framework, we would expect the amount of excess charge to be reduced significantly as compared to Fig. 6.

In Fig. 7 we show time-dependent currents for the same double barrier as in Fig. 5 for two different ways of applying the bias in the left lead: in one case the constant bias U_0 is switched on suddenly at $t=0$ (as in Fig. 5); in the other case the constant U_0 is achieved with a smooth switching $U(t) = U_0 \sin^2(\omega t)$ for $0 < t < \pi/(2\omega)$. As expected and in agreement with the results of Ref. 15, the same steady state is achieved after the initial transient time. However, the transient current clearly depends on how the bias is switched on.

In the final example we address the simulation of ac transport. We computed the current for a single square potential barrier with $V(x) = 0.6$ a.u. for $|x| < 6$ a.u. and zero otherwise. Here we applied a time-dependent bias of the form $U_L(t) = U_0 \sin(\omega t)$ to the left lead. The right lead remains on zero bias. The numerical parameters are Fermi energy $\varepsilon_F = 0.5$ a.u. and device region from $x = -6$ to $x = +6$ a.u. with lattice spacing $\Delta x = 0.03$ a.u. The number of k points is 100 and the time step is $\Delta t = 10^{-2}$ a.u. In Fig. 8 we plot the cur-

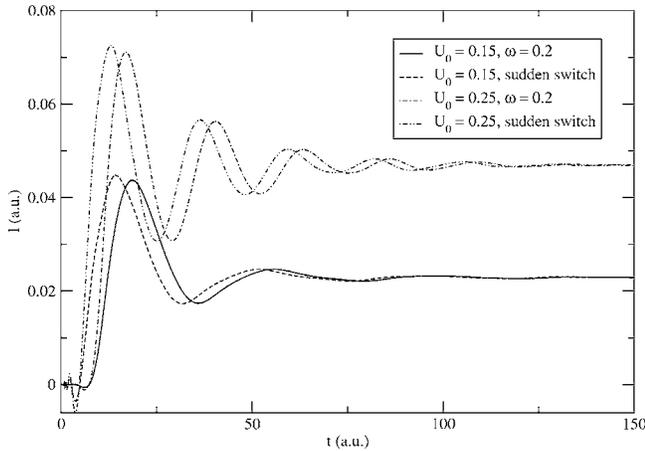


FIG. 7. The time evolution of the current for a double square potential barrier when the bias is switched on in two different manners: in one case, the bias U_0 is suddenly switched on at $t=0$ while in the other case the same bias is achieved with a smooth switching $U(t)=U_0 \sin^2(\omega t)$ for $0 < t < \pi/(2\omega)$. The parameters for the double barrier and the other numerical parameters are the same as the ones used in Fig. 5. U_0 and ω given in atomic units.

rent at $x=0$ as a function of time for different values of the parameter $U_0=0.1, 0.2, 0.3$ a.u. The frequency was chosen as $\omega=1.0$ a.u. in both cases. Again, as for the dc calculation discussed above, we get a transient that overshoots the average current flowing through the constriction; again, this excess current is larger the higher the applied voltage. Also, after the transient we obtain a current through the system with the same period as the applied bias. Note, however, that (especially for the large bias) the current is not a simple harmonic like the applied ac bias.

VII. SUMMARY AND OUTLOOK

In the present work we have presented a formally rigorous approach towards the description of charge transport using

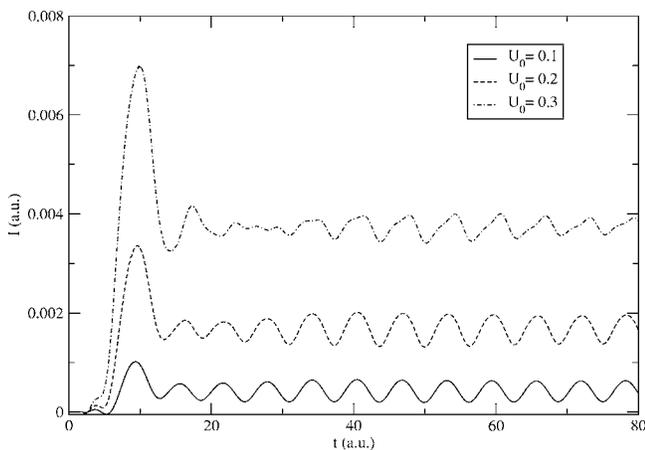


FIG. 8. The time evolution of the current for a square potential barrier in response to a time-dependent, harmonic bias in the left lead, $U_L(t)=U_0 \sin(\omega t)$ for different amplitudes U_0 (values in a.u.) and frequency $\omega=1.0$ a.u. The potential is given by $V(x)=0.6$ a.u. for $|x| \leq 6.0$ a.u. and zero otherwise. The propagation region extends from $x=-6$ to $x=+6$ a.u. The Fermi energy of the initial state is $\epsilon_F=0.5$ a.u. The current at $x=0$ is shown.

an open-boundary scheme within TDDFT. We have implemented a specific time-propagation scheme that incorporates transparent boundaries at the device/lead interface in a natural way. In order to have a clear definition of a device region we assumed that an applied bias can be described by adding a spatially constant potential shift in the macroscopic part of the leads. This implies an effective “metallic screening” of the constriction. The screening limits the spatial extent of the induced density created by the bias potential or the external field to the central region. Our treatment can be further refined to include dynamical screening deep inside the electrodes on the level of linear response, which might be of importance for the initial transient. Our time-dependent scheme allows us to extract both ac and dc I/V device characteristics and it is ideally suited to describe external field (photon) assisted processes.

In order to illustrate the performance of the method, we have implemented it for one-dimensional models and we have shown (i) how to extract the proper initial extended states to be propagated, (ii) how to incorporate perfect transparent boundaries for the time propagation, and (iii) that a steady-state current is always reached upon application of a dc bias. The transient process occurs on a time scale much shorter than the relaxation time due to electron-phonon interaction. The important observation that a steady state is reached without dissipation whenever large leads are coupled to a small device region was made in several explicitly time-dependent approaches^{15,29,37,38} and is, after all, also the basic idea behind the Landauer-Büttiker formulation. In the case of systems without any source of dissipation it is known that the steady current is independent of the history of the process.¹⁵ We have explicitly demonstrated this history independence for two different switching processes of the external bias. However, if we allow for dissipation either through electron-electron or electron-phonon interactions, the current versus voltage characteristics might depend on the history. For instance, hysteresis loops due to different transient electronic/geometrical device configurations are possible. This effect will be more dramatic in the case of ac-driving fields of high frequencies where the system may not have enough time to respond to the perturbation. We have also shown that (iv) a periodic alternating current is reached upon perturbation with a monochromatic field.

Previous work on time-dependent quantum transport mainly uses the idea of Caroli.^{9,10} This approach is at the core of the Landauer derivation and has the problem of using different chemical potentials in different parts of the system. This implies that the initial state is not a ground state of the entire, contacted system. Furthermore, the time-dependent perturbation is a tunneling Hamiltonian that is nonlocal in space. Therefore, it cannot be combined with TDDFT since there the time-dependent potential is local.

Here, we have used the scheme of Cini,¹⁴ which can be combined with TDDFT: We start the calculation from a well-defined thermodynamic equilibrium configuration, therefore the scheme is thermodynamically consistent. Then we apply an external potential that in general is time dependent. By virtue of the Runge-Gross theorem, the time evolution of this quantum system is completely determined by the knowledge of the time-dependent density. In the steady-state regime the

occupation of left- and right-moving carriers is dictated by the time-dependent Schrödinger equation.

At the level of linear response, other thermodynamically consistent schemes have been put forward using the Kubo formalism,³⁹ which both overcome the problem of having two chemical potentials. Bokes and Godby⁴⁰ have developed a linear response formalism working directly with the external and/or total electric field. Kamenev and Kohn⁴¹ have considered a closed system (ring) where the current in the system results from an external driving vector potential. They have shown that it is possible to recover the Landauer result, i.e., the universal quantum of conductance $2e^2/h$ independent of the length and material. Since the Kamenev-Kohn approach uses a vector potential rather than a scalar potential, time-dependent current DFT rather than TDDFT would be the natural density-functional extension.

Most theoretical approaches to transport in molecular electronic devices adopt open boundary conditions and assume that transport is ballistic, i.e., under steady-state conditions inelastic collisions are absent from a molecular structure and its contacts.^{4-6,33} Dissipation occurs only in the idealized macroscopic reservoirs connected by leads to the molecular device. This central role of the reservoirs in the process of dissipation is a valid approximation, particularly when the applied bias is small and a device operates in a linear regime. When inelastic scattering dominates, this picture is not applicable. In particular, experiments^{47,48} indicate that inelastic scattering with lattice vibrations is present at sufficiently large bias, causing local heating⁴²⁻⁴⁶ of contacts and molecular devices. Energy transfer to the lattice may also cause atomic migration and result in dramatic changes in the device characteristics (also they may give phonon replica structure in the measured conductance). The modeling of a many-electron system out of equilibrium coupled to lattice vibrations is a real theoretical challenge.⁴⁹

Electron correlations are also important in molecular conductors, for example, Coulomb blockade effects dominate the transport in quantum dots. Short-range electron correlations seems to be relevant in order to get a quantitative description of I/V characteristics in molecular constrictions.^{21,50,51} In particular, it is commonly assumed that the energy scales for electron-electron and electron-phonon interactions are different and could be treated separately. However, the metallic screening of the electrodes considerably reduces the Coulomb-charging energy (from eV to meV scale). In this regime the energy scales for the two interactions merge and they need to be treated on the same footing, posing some additional theoretical challenge.

Other approaches put forward in the literature directly look for a homogeneous current-carrying state either based on a maximum entropy principle⁵² or by imposing the current through Lagrange multipliers.⁵³ In those approaches it is implicitly assumed that the origin of the homogeneous current is independent whether it is introduced by reservoirs or by external fields. This is indeed the case for independent electrons, but once dissipation is built in the system might exhibit a dependence on the history of the applied bias (e.g., possible appearance of hysteresis loops⁴⁹ in the current versus voltage characteristics).

It is clear that the quality of the TDDFT functionals is of crucial importance. In particular, exchange and correlation

functionals for the nonequilibrium situation are required. Time-dependent linear response theory for dc steady state has been implemented in Ref. 38 within TDLDA assuming jellium-like electrodes (mimicked by complex absorbing/emitting potentials). It has been shown that the dc conductance changes considerably from the standard Landauer value. Therefore, a systematic study of the TDDFT functionals themselves is needed. A step beyond standard adiabatic approximations and exchange-only potentials is to resort to many-body schemes as recently done for the characterization of optical properties of semiconductors and insulators.⁵⁴ The derivative discontinuities recently discovered⁵⁵ in the case of ionization processes may play an important role for transport processes as well. Another path is to explore explicitly current-dependent approximations of the exchange-correlation potential.⁵⁶⁻⁵⁸

An appealing feature of the present approach is that electron-electron and electron-ion correlations and dissipation would, in principle, be described correctly in two-component TDDFT.⁵⁹

At present we are implementing our propagation scheme for real 3D systems³⁶ within the real-space real-time TDDFT code, OCTOPUS.⁶⁰ We are also exploring the possibility of a semiclassical description of the electron-ion coupling in order to avoid the complexity of multicomponent DFT and the problems related to mixed quantum classical approaches (i.e., Ehrenfest dynamics) as they fail to describe the long-term inelastic electron-phonon scattering correctly.^{49,61,62}

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APPENDIX: CONTINUED MATRIX FRACTIONS

Let us consider the infinite tridiagonal block matrix

$$M_0 = \begin{bmatrix} A_0 & B_1 & 0 & \dots \\ B_1 & A_1 & B_2 & \dots \\ 0 & B_2 & A_2 & \dots \\ \dots & \dots & \dots & \dots \end{bmatrix}, \quad (\text{A1})$$

where A_i and B_i are $N \times N$ matrices (the argument works even for matrices of different matching dimensions). We write the inverse matrix M_0^{-1} as

$$M_0^{-1} = \begin{bmatrix} Q_{00} & \tilde{Q}_{01} \\ \tilde{Q}_{10} & Q_1 \end{bmatrix}, \quad (\text{A2})$$

where Q_{00} is the first $N \times N$ block of M_0^{-1} . It is now convenient to introduce the matrix M_n obtained by dropping the first nN lines and nN columns of M_0 . Then, in terms of the

rectangular matrix $\tilde{B}_n = [B_n, 0, 0, \dots]$, we have

$$\begin{aligned} \begin{array}{c} 1 \\ \left[\begin{array}{cc} A_0 & \tilde{B}_1 \\ \tilde{B}_1^T & M_1 \end{array} \right] \end{array} &= \begin{array}{c} 1 \\ \left[\begin{array}{cc} A_0 & 0 \\ 0 & M_1 \end{array} \right] \end{array} \\ &- \begin{array}{c} 1 \\ \left[\begin{array}{cc} A_0 & 0 \\ 0 & M_1 \end{array} \right] \end{array} \begin{array}{c} \left[\begin{array}{cc} 0 & \tilde{B}_1 \\ \tilde{B}_1^T & 0 \end{array} \right] \end{array} \begin{array}{c} 1 \\ \left[\begin{array}{cc} A_0 & \tilde{B}_1 \\ \tilde{B}_1^T & M_1 \end{array} \right] \end{array}. \end{aligned} \quad (\text{A3})$$

From the above Dyson-like equation it is straightforward to obtain $Q_{00} = A_0^{-1} - A_0^{-1} \tilde{B}_1 \tilde{Q}_{10}$ and $\tilde{Q}_{10} = -M_1^{-1} \tilde{B}_1^T Q_{00}$. Solving for Q_{00} ,

$$Q_{00} = \frac{1}{A_0 - \tilde{B}_1 M_1^{-1} \tilde{B}_1^T}. \quad (\text{A4})$$

One can now proceed along similar lines. We define

$$M_1^{-1} = \begin{bmatrix} Q_{11} & \tilde{Q}_{12} \\ \tilde{Q}_{21} & Q_2 \end{bmatrix}, \quad (\text{A5})$$

where Q_{11} is the first $N \times N$ block of M_1^{-1} . From the corresponding Dyson equation one finds $Q_{11} = A_1^{-1} - A_1^{-1} \tilde{B}_2 \tilde{Q}_{21}$ and $\tilde{Q}_{21} = -M_2^{-1} \tilde{B}_2^T Q_{11}$. Solving for Q_{11}

$$Q_{11} = \frac{1}{A_1 - \tilde{B}_2 M_2^{-1} \tilde{B}_2^T} \quad (\text{A6})$$

and substituting this result back in Q_{00} yields

$$Q_{00} = \frac{1}{A_0 - B_1 \frac{1}{A_1 - \tilde{B}_2 M_2^{-1} \tilde{B}_2^T} B_1}. \quad (\text{A7})$$

Repeating the same steps we end up with the continued matrix fraction

$$Q_{00} = \frac{1}{A_0 - B_1 \frac{1}{A_1 - B_2 \frac{1}{A_2 - B_3 \frac{1}{A_3 - B_4 \frac{1}{A_4 \cdots} B_4} B_3} B_2} B_1}. \quad (\text{A8})$$

¹M. A. Reed, C. Zhou, C. J. Muller, T. P. Burgin, and J. M. Tour, *Science* **278**, 252 (1997).

²S. J. Tans, M. H. Devoret, H. Dai, A. Thess, R. E. Smalley, L. J. Geerligs, and C. Dekker, *Nature (London)* **386**, 474 (1997).

³J. Jortner and M. Ratner, *Molecular Electronics* (Blackwell Science, Oxford, 1997).

⁴For a recent overview of the field see *Lecture Notes in Physics, Introducing Molecular Electronics* (Springer Verlag, New York, 2005), and references therein. (<http://www-mcg.uni-regensburg.de/pages/admol/book/>)

⁵R. Landauer, *IBM J. Res. Dev.* **1**, 233 (1957).

⁶M. Büttiker, *Phys. Rev. Lett.* **57**, 1761 (1986).

⁷L. P. Kadanoff and G. Baym, *Quantum Statistical Mechanics* (Benjamin, New York, 1962).

⁸L. V. Keldysh, *Sov. Phys. JETP* **20**, 1018 (1965).

⁹C. Caroli, R. Combescot, P. Nozières, and D. Saint-James, *J. Phys. C* **4**, 916 (1971).

¹⁰C. Caroli, R. Combescot, D. Lederer, P. Nozières, and D. Saint-James, *J. Phys. C* **4**, 2598 (1971).

¹¹A.-P. Jauho, N. S. Wingreen, and Y. Meir, *Phys. Rev. B* **50**, 5528 (1994).

¹²Y. Meir and N. S. Wingreen, *Phys. Rev. Lett.* **68**, 2512 (1992).

¹³H. Haug and A.-P. Jauho, *Quantum Kinetics in Transport and Optics of Semiconductors* (Springer, Berlin, 1998).

¹⁴M. Cini, *Phys. Rev. B* **22**, 5887 (1980).

¹⁵G. Stefanucci and C.-O. Almbladh, *Phys. Rev. B* **69**, 195318 (2004).

¹⁶N. D. Lang, *Phys. Rev. B* **52**, 5335 (1995).

¹⁷P. A. Derosa and J. M. Seminario, *J. Phys. Chem. B* **105**, 471 (2001).

¹⁸M. Brandbyge, J.-L. Mozos, P. Ordejón, J. Taylor, and K. Stokbro, *Phys. Rev. B* **65**, 165401 (2002).

¹⁹Y. Xue, S. Datta, and M. A. Ratner, *Chem. Phys.* **281**, 151 (2002).

²⁰A. Calzolari, N. Marzari, I. Souza, and M. B. Nardelli, *Phys. Rev. B* **69**, 035108 (2004).

²¹F. Evers, F. Weigend, and M. Koentopp, *Phys. Rev. B* **69**, 235411 (2004).

²²P. S. Krstić, D. J. Dean, X. G. Zhang, D. Keffer, Y. S. Leng, P. T. Cummings, and J. C. Wells, *Comput. Mater. Sci.* **28**, 321 (2003).

²³E. Runge and E. K. U. Gross, *Phys. Rev. Lett.* **52**, 997 (1984).

²⁴M. Petersilka, U. J. Gossmann, and E. K. U. Gross, *Phys. Rev. Lett.* **76**, 1212 (1996).

²⁵G. Stefanucci and C.-O. Almbladh, *Europhys. Lett.* **67**, 14 (2004).

²⁶C. Moyer, *Am. J. Phys.* **72**, 351 (2004) and references therein.

²⁷P. Hohenberg and W. Kohn, *Phys. Rev.* **136**, B864 (1964).

²⁸The original proof of the Runge-Gross theorem referred to finite systems. The proof, however, is not restricted to finite systems as long as the perturbing fields have finite range. The precise conditions of validity were laid down in E. K. U. Gross, and W. Kohn, *Adv. Quantum Chem.* **21**, 55 (1990). The particular case of periodic solids was discussed in detail in N. T. Maitra, I.

- Souza and K. Burke, Phys. Rev. B **68**, 045109 (2003).
- ²⁹M. Di Ventra and T. N. Todorov, J. Phys.: Condens. Matter **16**, 8025 (2004).
- ³⁰J. R. Hellums and W. R. Frensley, Phys. Rev. B **49**, 2904 (1994).
- ³¹We should remark here that in a TDDFT description of a fully interacting many-electron system other memory terms should appear due to dynamical exchange-correlation effects and initial state-dependence. [See J. F. Dobson, M. J. Büchner, and E. K. U. Gross, Phys. Rev. Lett. **79**, 1905 (1997) and R. van Leeuwen, Phys. Rev. Lett. **82**, 3863 (1999) for a procedure to build this dependence in the potential.] Most density-based functionals used at present rely on the adiabatic approximation, therefore ignoring the functional dependence on past time-dependent densities (Ref. 32).
- ³²N. T. Maitra, K. Burke, and C. Woodward, Phys. Rev. Lett. **89**, 023002 (2002).
- ³³S. Datta, *Electronic Transport in Mesoscopic Systems* (Cambridge University Press, Cambridge, 1995).
- ³⁴D. S. Fisher and P. A. Lee, Phys. Rev. B **23**, R6851 (1981).
- ³⁵Higher order propagation schemes could be developed following the ideas discussed here for the modification of the Cayley method. Such schemes could also handle self-consistent and time-dependent Hamiltonians as the ones appearing in standard TDDFT. See A. Castro, M. A. L. Marques, and A. Rubio, J. Chem. Phys. **121**, 3425 (2004) and references therein.
- ³⁶L. Wirtz, G. Stefanucci, C.-O. Almbladh, S. Kurth, H. Appel, E. K. U. Gross, and A. Rubio (unpublished).
- ³⁷N. Bushong, N. Sai, and M. Di Ventra, cond-mat/0504538
- ³⁸R. Baer, T. Seideman, S. Ilani, and D. Neuhauser, J. Chem. Phys. **120**, 3387 (2004).
- ³⁹R. Kubo, M. Toda, and N. Hashitsume, *Statistical Physics II: Nonequilibrium Statistical Mechanics* (Springer, Berlin, 1991).
- ⁴⁰P. Bokes and R. W. Godby, Phys. Rev. B **69**, 245420 (2004).
- ⁴¹A. Kamenev and W. Kohn, Phys. Rev. B **63**, 155304 (2001).
- ⁴²A. P. Horsfield, D. R. Bowler, and A. J. Fisher, J. Phys.: Condens. Matter **16**, L65 (2004).
- ⁴³Y. C. Chen, M. Zwolak, and M. Di Ventra, Nano Lett. **3**, 1691 (2003).
- ⁴⁴M. J. Montgomery, T. N. Todorov, and A. P. Sutton, J. Phys.: Condens. Matter **14**, 5377 (2002).
- ⁴⁵M. J. Montgomery and T. N. Todorov, J. Phys.: Condens. Matter **15**, 8781 (2003).
- ⁴⁶A. Troisi, M. A. Ratner, and A. Nitzan, J. Chem. Phys. **118**, 6072 (2003).
- ⁴⁷J. G. Kushmerick, J. Lazorcik, C. H. Patterson, R. Shashidhar, D. S. Seferos, and G. C. Bazan, Nano Lett. **4**, 639 (2004).
- ⁴⁸R. H. M. Smit, C. Untiedt, and J. M. van Ruitenbeek, Nanotechnology **15**, S472 (2004).
- ⁴⁹A different approach based on the time-dependent evolution of a Liouville-master equation in the presence of dissipative environment has been put forward by R. Gebauer and R. Car, Phys. Rev. Lett. **93**, 160404 (2004); Phys. Rev. B **70**, 125324 (2004); K. Burke, R. Car, and R. Gebauer, Phys. Rev. Lett. **94**, 146803 (2005).
- ⁵⁰P. Delaney and J. C. Greer, Phys. Rev. Lett. **93**, 036805 (2004).
- ⁵¹A. Ferretti, A. Calzolari, R. Di Felice, F. Manghi, M. J. Caldas, M. B. Nardelli, and E. Molinari, Phys. Rev. Lett. **94**, 116802 (2005).
- ⁵²P. Bokes and R. W. Godby, Phys. Rev. B **68**, 125414 (2003).
- ⁵³D. S. Kosov, J. Chem. Phys. **120**, 7165 (2004).
- ⁵⁴A. Marini, R. Del Sole, and A. Rubio, Phys. Rev. Lett. **91**, 256402 (2003); L. Reining, V. Olevano, A. Rubio, and G. Onida, *ibid.* **88**, 066404 (2002); I. V. Tokatly and O. Pankratov, *ibid.* **86**, 2078 (2001).
- ⁵⁵M. Lein and S. Kümmel, Phys. Rev. Lett. **94**, 143003 (2005).
- ⁵⁶C. A. Ullrich and G. Vignale, Phys. Rev. B **65**, 245102 (2002) and references therein.
- ⁵⁷I. V. Tokatly and O. Pankratov, Phys. Rev. B **67**, 201103(R) (2003).
- ⁵⁸N. Sai, M. Zwolak, G. Vignale, and M. Di Ventra, cond-mat/0411098.
- ⁵⁹T. Kreibich, "Multicomponent density-functional theory for molecules in strong laser fields," Ph.D. thesis, Universität Würzburg, Shaker-Verlag (2000); T. Kreibich, R. van Leeuwen, and E. K. U. Gross, Chem. Phys. **304**, 183 (2004) and references therein.
- ⁶⁰M. A. L. Marques, A. Castro, G. F. Bertsch, and A. Rubio, Comput. Phys. Commun. **151**, 60 (2003). The code OCTOPUS is available at <http://www.tddft.org/programs/octopus>.
- ⁶¹A. Castro, X. Lopez, M. A. L. Marques, A. Rubio, E. K. U. Gross (unpublished).
- ⁶²A. P. Horsfield, D. R. Bowler, A. J. Fisher, T. N. Todorov, and M. J. Montgomery, J. Phys.: Condens. Matter **16**, 3609 (2004).