Fermi-edge problem in the presence of AC electric field

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Abstract – In this paper, we study a non-equilibrium Fermi-edge problem where the system under investigation is a single electron reservoir put under an AC electric field. We show that the electron Green’s function and other correlation functions in the problem can be solved and expressed exactly in terms of a well-defined integral. The qualitative behaviors of the solution is studied and compared with the situation where the impurity is coupled to more than one reservoir at different chemical potentials.

The Fermi-edge problem in non-equilibrium situation is perhaps one of the most simple but non-trivial problems in non-equilibrium quantum many-body physics. It is the generalization of the Nozières-De Dominicis X-ray edge problem [1] to non-equilibrium situations where the systems may be i) connected to \( N > 1 \) reservoirs at different chemical potentials [2], ii) put under a time-dependent electromagnetic field, or iii) put under other non-equilibrium configurations. The problem concerns the reaction of a non-equilibrium electron gas to the sudden turning on of a scattering potential. A simple realization of the problem is a small quantum dot coupled to electron reservoirs as shown in fig. 1. The small quantum dot can be treated as a two-level system, and by changing the gate voltage applied to the quantum dot, the electron occupancy of the dot can be changed, resulting in a change of the local dot potential felt by electrons in the reservoirs [3–5]. Indeed, Fermi-edge singularity behavior was observed experimentally in a quantum dot system at energy range \( \sim 1 \text{ meV} \) [6]. The problem is in principle a one-particle problem, where the many-body effects enter only through Fermi statistics. The problem has been studied carefully in those cases where the system is connected to reservoirs at different chemical potentials [3,7–9]. It is now understood that in this case the problem is equivalent to solving an \( N \times N \) matrix Riemann-Hilbert (RH) boundary value problem [7,8] of which a general exact solution is unavailable and only the long-time asymptotic behavior of the solution can be extracted [2,7,8].

Fig. 1: (Color online) (a) \( N \) reservoirs at different chemical potentials coupled to a quantum dot. (b) A single reservoir coupled to a quantum dot. The system is put under an AC electric field. (c) Dissipation caused by chemical potential differences rounds off the Fermi-edge singularities. (d) Sidebands will appear for electrons in a single reservoir under an AC field.

In this paper, we consider a different situation where the system has only one electron reservoir put under an AC electric field generated by, for example, a microwave radiation. Going to the center-of-mass (CM) frame, it can be shown that the electric field can be eliminated [10] and that the system becomes an “equilibrium” system coupled to a time-dependent (periodic) impurity scattering potential. The corresponding Fermi-edge problem concerns the reaction of this “equilibrium” electron gas to the sudden turning on of a time-dependent scattering potential. The problem provides a rare example of a non-trivial non-equilibrium quantum many-body problem
that can be solved exactly and is testable. We find that the equation of motion for the electron Green’s function is a Carleman-type integral equation of which the solution can be expressed in terms of a well-defined integral. In this paper, the qualitative behaviors of the solution are analyzed and compared with the situation where the impurity potential is static, but the system is connected to reservoirs at different chemical potentials. 

The system that we consider is described by the time-dependent one-particle Hamiltonian \( H = H_0(t) + V(t) \), where

\[
H_0(t) = -\sum_i \frac{\hbar^2}{2m} \nabla_i^2 - e \sum_i \vec{r}_i \cdot \vec{E}(t),
\]

(1a)

where the first term represents the kinetic energy of electrons, \( \vec{E}(t) \) represents the AC electric field, \( \vec{r}_i \) is the position of the \( i \)-th electron and

\[
V(t) = \theta(t_f - t) \theta(t - t_i) \sum_i U(\vec{r}_i)
\]

(1b)

describes a scattering potential \( U(\vec{r}) \) located at the origin turning on (off) at time \( t = t_i(t_f) \). We have neglected spin as in usual practice.

In the absence of \( V(t) \), the electric field couples only to the CM coordinate \( \vec{R} = 1/N \sum_i \vec{r}_i \) of the whole electron fluid and can be eliminated by performing a coordinate transformation to CM frame of the system [10]. After a gauge transformation, the Hamiltonian in CM frame becomes

\[
H_{0(CM)} = \sum_i -\frac{\hbar^2}{2m} \nabla_i^2,
\]

(2a)

where \( \vec{r}_i' = \vec{r}_i - \vec{R}(t) \), \( \vec{R}(t) \) is the CM trajectory determined by the classical equation of motion \( m \vec{R} = e \vec{E}(t) \). In the transformed coordinate, \( V(t) \) becomes

\[
V_{CM}(t) = \theta(t_f - t) \theta(t - t_i) \sum_i U(\vec{r}_i' + \vec{R}(t)),
\]

(2b)

where the original static impurity becomes a moving impurity in the CM frame with corresponding time-dependent scattering potential \( U_{CM}(\vec{r},t) = U(\vec{r} + \vec{R}(t)) \).

As in equilibrium situation, we shall assume that the scattering events can be decomposed into separate scattering channels, and keep only one channel in our study. In this case, we may replace the scattering potential by \( U_{CM}(t) \sim U_0(t) \delta(\vec{r}_i') \), where \( U_0(t) \) is a periodic potential in time. With this approximation, the only difference between the equilibrium Fermi-edge problem and the present problem is the appearance of a time-dependent scattering potential \( U_0(t) \). In realistic situations, different scattering channels may be mixed by the traveling impurity. The qualitative behavior of the Fermi-edge singularity is not modified by this mixing [8] as long as the scattering potentials in different channels have the same periodicity.

Before presenting the solution, we first revisit the case of a static impurity coupled to two reservoirs at different chemical potentials \( \mu_1 \) and \( \mu_2 \). In this case, the chemical potential difference can be gauged away resulting in an effective time-dependent impurity scattering, \( V_{12}(t) \sim V_0 e^{-i\Delta\mu} (\Delta \mu = \mu_1 - \mu_2) \), that couples the two reservoirs. The long-time \( (t_f - t, t - t_i \gg \hbar/\Delta \mu) \) response of the electron gas to the suddenly switched on impurity potential is determined by two phase shifts associated with the two reservoirs. The phase shifts are complex when \( \Delta \mu \neq 0 \) and reflect the existence of dissipation in an out-of-equilibrium system. Associated with the two phase shifts are two Fermi-edge singularities which are rounded off by finite life-time effects associated with the imaginary part of the complex phase shifts [2,3,7,8]. For electrons in a single reservoir under a time-dependent electric field with period \( T = 2\pi/\omega_0 \), side-bands will appear in the electron spectral function at frequencies \( \omega \sim \mu + n \omega_0 \), where \( n \) is integer. An impurity will cause scattering between different side-bands. Will Fermi-edge singularities appear on all these side-bands and will the singularities be rounded off by finite life-time effect as in the many-reservoir situation? We note that both AC and DC transports are usually dissipative in condensed matter systems. Will there be other new effects? These are the questions to be asked in the present problem.

At zero temperature, the main challenge of the Fermi-edge problem is to evaluate the one-particle time-ordered Green’s function \( G(t,t') \) satisfying the Dyson equation,

\[
G(t,t') = g(t-t') + i \int_{t_i}^{t_f} g(t-t'')U_0(t'')G(t'',t') \, dt'',
\]

(3a)

where

\[
g(t-t') \sim -i\nu_0 \left( P \frac{1}{t-t'} + \pi \tan \theta \delta(t-t') \right)
\]

(3b)

is the unperturbed Green’s function determined by \( H_0 \). Green’s functions are defined in the CM coordinate in eq. (3). A similar equation can also be derived at finite temperature [9] with \( g \) replaced by its finite temperature extension. Since the Hamiltonian is quadratic, other correlation functions can be computed once \( G \) is determined. An important correlation function characterizing the Fermi-edge singularity is the propagator

\[
B(t_f,t_i) = e^{C(t_f,t_i)} \langle \psi_0 | S(t_f,t_i) | \psi_0 \rangle,
\]

where \( S(t_f,t_i) = T \exp(-i/\hbar \int_{t_i}^{t_f} U_0(t') \, dt') \) is the time evolution operator under the scattering potential \( U_0(t) \) and \( | \psi_0 \rangle \) is the ground state wavefunction of the system (in the CM frame) before \( U_0(t) \) is switched on. \( C(t_f,t_i) \) measures the overlap between \( | \psi_0 \rangle \) and the final state after \( U_0(t) \) is switched on (orthogonality catastrophe [11]) and is related to \( G \) by

\[
\lambda \frac{\partial C(t_f,t_i)}{\partial \lambda} = i \int_{t_i}^{t_f} \langle \lambda U_0(t) \rangle G^\lambda(t,t) \, dt,
\]

(4)
where $G^A$ is Green’s function determined from eq. (3) with $U_0(t) \rightarrow \lambda U_0(t)$ [1]. The electron Green’s function measuring the Fermi-edge singularity is [1]

$$K(t_f, t_i) = G(t \rightarrow t_f, t' \rightarrow t_i)e^{C(t_f, t')}.$$ 

To solve eq. (3), we introduce $\tilde{G}(t, t') = \nu_0 U_0(t)G(t, t')$. After some simple manipulation, we obtain

$$\alpha(t)\tilde{G}(t, t') = g(t - t') + P \int_{t_i}^{t_f} \frac{\tilde{G}(t'', t')}{t - t''} dt'', \quad (5)$$

where $\alpha(t) = \frac{1 - \pi \nu_0 \tan \theta_0 \tilde{U}_0(t)}{\nu_0 \lambda(t)}$. The equation is a Carleman-type equation with $t'$ as a dummy variable. The solution to this equation can be constructed by analyzing the analytic behavior of the equation on the complex plane [12] (essentially an RH problem) and the relevant solution to our problem is

$$\tilde{G}(t, t') = \frac{\sin \delta(t)}{\pi} \times \left[ \cos \delta(t)g(t - t') + \frac{e^{-\varphi(t)}}{\pi} P \int_{t_i}^{t_f} \frac{e^{-\varphi(t'')}}{t - t''} \cos \delta(t'') dt'' \right], \quad (6a)$$

where $\delta(t) = \tan^{-1}(\pi/\alpha(t))$ and

$$\varphi(t) = \text{Re} \Phi(t + i\epsilon) = -\frac{1}{\pi} P \int_{t_i}^{t_f} \frac{\delta(t')}{t - t'} dt'. \quad (6b)$$

Notice that the complex function $\Phi(z) = -\frac{1}{\pi} \int_{t_i}^{t_f} \frac{\delta(t')}{z - t'} dt'$ is an analytical function on the complex plane except the segment $[t_i, t_f]$. Using eq. (3b) for $g$ and the fact that

$$-\frac{1}{\pi} P \int_{t_i}^{t_f} \frac{1}{t - t'} e^{\varphi(t')} \sin \delta(t') dt' = e^{\varphi(t)} \cos \delta(t)$$

at time $t_i < t < t_f$ because the integral is just the Kramers-Kronig relation relating the real and imaginary parts of the analytic function $\Phi(t + i\epsilon) = e^{\Phi(t + i\epsilon)}$, we obtain after some algebra,

$$G(t, t') = (-i\nu_0)A_0(t) \times \left[ e^{-\varphi(t)}e^{\varphi(t')}A_0(t') \frac{1}{t - t'} + \cos \delta(t) \tan \theta_0 - \sin \delta(t) \right] \pi \delta(t - t') \quad (7)$$

for $t_i < t < t'$, where $A_0(t) = (1 + \tan \delta(t) \tan \theta_0)\cos \delta(t)$.

First we examine eq. (7) in the equilibrium situation $U_0(t) = 0$. In this case, $\delta(t) \rightarrow \delta$ becomes a constant and $\varphi(t) \rightarrow \frac{\pi}{2} \ln(t - t_i)$. It is easy to see that the ND result for the electron Green’s function [1] is recovered. The non-trivial result here is that the phase shift $\delta(t)$ becomes time-dependent because of the time-dependent potential $U_0(t)$. Notice that contrary to the case of a static impurity coupled to reservoirs at different chemical potentials where the relevant phase shifts are complex, the phase shift $\delta(t)$ here is always real.

It is straightforward to show that

$$\tilde{G}(t, t' \rightarrow t) = (-i\nu_0)\frac{1}{\pi} \sin \delta(t) A_0(t) \left( i\epsilon_c - \frac{\partial \varphi(t)}{\partial t} \right),$$

where $\epsilon \sim \lim_{t \rightarrow t'} \frac{1}{\tau}$ is a high-energy cutoff below which the approximate expression (3b) for $g_0$ is valid [1]. Putting this back into eq. (4) and using the identity $\text{d}y/\lambda = \delta/(\text{d} \sin \lambda)$, we obtain after some simple algebra

$$C(t_f, t_i) = \int_{t_i}^{t_f} dt \int_{0}^{\delta(t)} \frac{\delta(t) \text{d} \delta(t)}{\pi} (\epsilon \epsilon_c - \partial \varphi(t)/\partial t), \quad (8)$$

In the equilibrium case $\delta(t) = \delta$, the first term $\epsilon \int_{t_i}^{t_f} \text{d} \delta(t)$ in eq. (8) represents a self-energy correction to ground state energy from impurity scattering (Fumi’s theorem [13]). The second term introduces logarithmic corrections to $C(t_f, t_i)$ coming from suddenly switching on/off the scattering potentials at $t_i(t_f)$ (orthogonality catastrophe) [1,11]. Equation (8) represents a generalization of this result to time-dependent scattering potentials. We shall consider the situation where $\alpha(t)$ is periodic in time in the following. In this case, $\delta(t)$ is also periodic in time and we can write $\delta(t) = \delta_0 + \sum_{n \neq 0} \delta_n e^{i\omega_n t}$, where $\delta_n = \delta - n \delta(t)$ is real.

$\varphi(t)$ cannot be evaluated exactly in this case. However, it is easy to extract the short- and long-time behaviors of $\varphi(t)$. In the short-time limit $\epsilon \ll t - t_i, t_f - t \ll T$, the leading contribution is

$$\varphi(t) \sim \frac{\delta(t)}{\pi} \ln \left( \frac{t_f - t}{t_i - t} \right), \quad (9)$$

whereas in the opposite limit $t_i - t, t_f - t \gg T$, we obtain

$$\varphi(t) = \frac{\delta_0}{\pi} \ln \left( \frac{t - t_i}{t - t_f} \right) + \varphi_0(t), \quad (10a)$$

where $\varphi_0(t) = \varphi^\infty(t) + \varphi_c(t)$ is the contribution from $\delta_n \neq 0$ terms in the Fourier representation of $\delta(t)$.

$$\varphi^\infty(t) = i \sum_{n \neq 0} \delta_n \text{sgn}(n) e^{i\omega_n t} \text{ is the contribution in the limit } -t_i = t_f \rightarrow \infty \text{ and } \varphi_c(t) = -i \sum_{n \neq 0} \frac{\delta_n}{\pi} \left[ e^{i\omega_n t} Ei(n\omega_0(t_f - t)) - e^{-i\omega_n t} Ei(n\omega_0(t_i - t)) \right] \quad (10b)$$

is the correction from finite time cutoff $t_i(t_f)$. $Ei(x) = P \int_{-\infty}^{x} e^y/a dy$ is the exponential integral function. It is easy to see that $\varphi^\infty(t)$ is a real and periodic function in time. Using the asymptotic results $Ei(z) \sim \ln z$ for $|z| \ll 1$ and $Ei(z) \sim e^z/z$ for $|z| \gg 1$, we find that $\varphi_c(t(t_i)) \sim b(t(t_i)) + \frac{1}{\pi \sqrt{t_i - t_f}} \int_{t_i}^{t_f} \text{d} \delta(t')$ in the limit $t_f - t_i \gg T$, where $b(t)$ is a periodic function in time and the second term is small compared with the $\delta_0$ term in eq. (10a).
With these results, we can now evaluate Green’s function $G(t \rightarrow t_f, t' \rightarrow t_i)$ in the long- and short-time limits $t_f - t_i \gg T$ and $\epsilon_c^{-1} \ll t_f - t_i \ll T$. In the long-time limit, we obtain

$$G(t \rightarrow t_f, t' \rightarrow t_i) \rightarrow \frac{-iv_0}{t_f - t_i} A_-(t_f) \times (i\epsilon_c(t_f - t_i))^{2\hbar_0} A_+(t_i),$$

(11)

where $A_\pm(t) = A_0(t)e^{\mp \nu_0(t)}$ are periodic functions in time. The corresponding Green’s function in the short-time limit can be obtained by replacing $2\hbar_0$ by $\delta(t_i) + \delta(t_f) \sim 2\delta((t_i + t_f)/2)$ and $A_\pm(t)$ by $A_0(t)$ in eq. (11).

It is also straightforward to obtain in the long-time limit $t_f - t_i \gg T$

$$C(t_f, t_i) = i \int_{t_i}^{t_f} dt \frac{\delta(t)}{\pi} \epsilon_c - \frac{\delta_0^2}{\pi^2} \ln(i\epsilon_c(t_f - t_i)) \times \varphi(t_f) - \varphi(t_i),$$

(12)

where $\varphi(t)$ is a real and periodic function of time $t$ to leading order in $T/(t_f - t_i)$ coming from the $\varphi_n(t)$ term. The short-time behavior of $C(t_f, t_i)$ can be obtained by replacing $\delta_0^2$ by $\delta((t_f)^2 + \delta(t_i)^2)/2 \sim \delta((t_i + t_f)/2)^2$ and dropping the $\varphi(t)$ terms in eq. (12).

Combining these results, we find that the electron Green’s function $K = Ge^C$ can be written as

$$K(t_f, t_i) \sim K^0(t_f, t_i) (i\epsilon_c(t_f - t_i))^{\gamma((t_f + t_i)/2)}$$

(13)

in both the long- and short-time limits where $\gamma(t) \sim 2\delta(t)/\pi \sim \delta(t)^2/\pi^2$ in the short-time limit $\epsilon_c^{-1} \ll t_f - t_i \ll T$ and $\gamma(t) \rightarrow \gamma_0 = 2\hbar_0/\pi - \delta_0^2/\pi^2$ in the long-time limit $t_f - t_i \gg T$. $K^0(t, t') = A_-(t) P_{\frac{1}{2}} A_+(t')$, where $A_\pm(t)$ are periodic (but different) functions in time in the long- and short-time limits.

With eq. (13), we can evaluate the Fourier transformed Green’s function $K(\omega, \omega') = \int dt \int dt' e^{i\omega(t-t')} K(t, t') \theta(t - t')$ in the frequency regimes $\epsilon_c \gg |\omega + \omega'| \gg 2v_0$ and $|\omega + \omega'| \ll 2\omega_0$. In the low-frequency limit $|\omega + \omega'| \ll 2\omega_0$, we obtain

$$K(\omega, \omega') \sim \sum_n \delta(\omega - \omega' + n\omega_0) \alpha_n \left(\frac{\epsilon_c}{\omega + n\omega_0/2}\right)^{\gamma_0},$$

(14)

where $\alpha_n$ are coefficients which depend on the strength and polarization of the AC electric field.

In the high-frequency regime $2\epsilon_c \gg |\omega + \omega'| \gg 2\omega_0$, $K(\omega, \omega') \sim \int_{-\infty}^{\infty} dt_m e^{i(\omega - \omega') t_m} A_0(t_m)^2 \left(\frac{\epsilon_c}{\omega + n\omega_0}\right)^{\gamma_0}$. Evaluating the integral, we obtain

$$K(\omega, \omega') \sim \sum_n \delta(\omega - \omega' + n\omega_0) \alpha_n (\omega + \omega') \left(\frac{\epsilon_c}{\omega + n\omega_0/2}\right)^{\gamma_0},$$

(15)

where $\gamma_0 \sim (\gamma(t_m))$ is the critical exponent in the corresponding equilibrium Fermi-edge problem. $\alpha_n(\omega + \omega')$ are constants up to logarithmic correction factors. The Fermi-edge singularity in the high-frequency regime $|\omega + \omega'| \gg 2v_0$ is insensitive to the AC electric field. Notice that, in general, $\alpha_n \neq \alpha_n$ and $\gamma_0 \neq \gamma_s$.

Equations (14) and (15) predict that the Fermi-edge spectrum $ReK(\omega, \omega') = \sum_n \delta(\omega - \omega' + n\omega_0) S_n(\omega)$ develops side-bands as a result of the time-periodic potential. The spectral functions $S_n(\omega)$ develop power law singularities at $\omega \rightarrow -n\omega_0/2$ characterized by an $n$-independent Fermi-edge exponent $\gamma_0$ for $\omega + n\omega_0/2 \ll \omega_0$. The Fermi-edge singularities are not rounded off by finite life-time effects as in the case of systems with reservoirs at different chemical potentials. This is non-trivial since dissipation usually exists in both DC and AC transports in many-reservoir problems [2, 14]. In fact, it was shown in ref. [14] that dissipation exists in a system of quantum dot coupled to external reservoirs when the system is driven by a frequency $f \ll W$, where $W$ is the bandwidth of the quantum dot (quantum pump problem). The finite quantum dot bandwidth provides a natural mechanism for dissipation. Our system consists of only one single impurity corresponds roughly to the opposite limit $f \gg W \rightarrow 0$ of ref. [14], where a natural mechanism for dissipation does not exist. The Fermi-edge singularity at the high-frequency regime $\epsilon_c \gg \omega \gg \omega_0$ is found to be characterized by an envelope function covering all side-bands with an exponent $\gamma_s$ which is insensitive to AC field and is different from the low-frequency exponent $\gamma_0$ (see fig. 2). These predicted features can be tested in mesoscopic systems under AC electric fields. In summary, we have analyzed in this paper a non-equilibrium Fermi-edge problem where the system under consideration is driven out-of-equilibrium by a time-dependent (AC) electric field. We showed that the system can be mapped into an equilibrium system with a traveling impurity or an impurity with a time-dependent scattering potential. We found that the problem can be solved exactly, and the corresponding Fermi-edge singularity behavior was very different from the case of a static impurity coupled to many reservoirs at different chemical potentials. Our solution provides a rare example.

\begin{figure}[h]
\centering
\includegraphics[width=0.8\textwidth]{fig2.png}
\caption{Spectral function in the presence of AC field.}
\end{figure}
of non-equilibrium quantum many-body problem which can be solved exactly.

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