Dissipation of Energy of an AC Electric Field in a Semi-Infinite Electron Plasma with Diffusive Boundary Conditions

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Abstract—Let us consider the electron plasma response with an arbitrary degree of degeneracy to an external ac electric field. Surface absorption of the energy of an electric field is calculated.

Index Terms—the Vlasov–Boltzmann equation, energy absorption of an electric field

I. INTRODUCTION

The character of electric field screening near the surface of a conductor is critically important for different problems of surface physics [1]–[2], in particular, the problem of propagation of plasma oscillations [3]–[4].

Here, we have obtained an analytical solution to the problem on the behavior of a semi-infinite plasma with an arbitrary degree of electron gas degeneracy in an external ac electric field perpendicular to the plasma surface. Such a situation takes place, e.g., when analyzing a solid-state semiconductor plasma. We use the Vlasov–Boltzmann kinetic equation with the Bhatnagar–Gross–Krook (BGK) collision equation with the locally equilibrium Fermi–Dirac distribution function, $f_\text{eq}$ is the unperturbed Fermi–Dirac distribution function, $f_\text{eq}(v, \mu_0) = f_\text{FD}(v, \mu_0)$, $\text{FD}$ is the Fermi–Dirac distribution function, $\text{FD}$ is the electron distribution function; $\nu$ is the electron scattering frequency; $s$ is the particle spin ($s = 1/2$ for electrons); $k$ is the Boltzmann constant; $T$ is the plasma temperature, which is assumed to be constant; and $E(x, t)$ is the electric field in plasma.

Let us consider the condition of diffusive reflection of electrons from the boundary of a semi-infinite plasma: $f(x = 0, v, t) = f_\text{eq}(x = 0, v, t)$ at $v_x > 0$, $e(0) = 1$, $e(+\infty) < +\infty$. The external electric field on the plasma surface is perpendicular to the plasma boundary and varies in time as $E_{\text{ext}}(t) = E_0 e^{-i\omega t}(1, 0, 0)$.

The corresponding self-consistent electric field in plasma has the form $E(x, t) = E(x) e^{-i\omega t}(1, 0, 0)$.

We assume that the external field is sufficiently weak, so that the linear approximation is applicable. Equations (1) and (2) can be linearized with respect to the absolute Fermi–Dirac distribution function $f_0$: $f_\text{eq}(x, P, t) = f_0(P, \alpha) + g(P, \alpha)e^{-i\omega t}$, where $f_0(P, \alpha) = f_\text{FD}(P, \alpha) = (1 + e^{\mu_0 - \alpha})^{-1}$, $g(P, \alpha) = e^{\mu_0 - \alpha} / (1 + e^{\mu_0 - \alpha})^2$. $P = p / p_T = v / v_T$. Here $v_T$ is the electron thermal velocity given by $v_T = \sqrt{2kT/m}$ and $\mu = \mu / kT$ is the reduced chemical potential. The change of the chemical potential is considered to be a small parameter so that representation $\alpha(x, t) = \mu + \delta \alpha(x)e^{-i\omega t}$ is possible. We linearize the electron distribution function $f_\text{eq}(x, P, \alpha) = f_0(P, \alpha) + g(P, \alpha)h(x, P_x)e^{-i\omega t}$, where $h(x, P_x)$ is a new unknown function and $h(x, P_x) \sim E$.

As a result, we get a system containing new unknown functions and dimensionless variables. The detailed solution is given in [6]. The solution is based on the method of separation of variables, is reduced to obtaining the dispersion function.
and search eigenfunctions by which we can decompose the resulting analytical solution. Dispersion function determines the range of solutions to the problem

\[ \Lambda(z) = 1 - \frac{1}{w_0} - \frac{z^2 - \eta_1^2}{\eta_0^2} \lambda_0(z, \alpha), \]

\[ \lambda_0(z, \alpha) = 1 + z \int_{-\infty}^{+\infty} \frac{k(\mu, \alpha) d\mu}{\mu - z}. \]

Constants \( w_0, \eta_1^2 \) and function \( k(\eta, \alpha) \) have forms

\[ f_0(\eta, \alpha) = (1 + \exp(\eta^2 - \alpha))^{-1}, \quad k(\eta, \alpha) = \frac{f_0(\eta, \alpha)}{2s_0(\alpha)} \]

\[ s_0(\alpha) = \int_{0}^{+\infty} f_0(t, \alpha) dt, \quad s_2(\alpha) = \int_{0}^{+\infty} t^2 f_0(t, \alpha) dt \]

\[ w_0 = 1 - \frac{i\omega}{\nu}, \quad \eta_1^2 = \frac{\nu^2 s_2(\alpha)}{\nu^2 s_0(\alpha)}. \]

As a result of the solution, the induced electromagnetic field is represented as the sum of three terms corresponding to the expansion in the spectrum of the dispersion function. In general, the structure of an electric field arising in a plasma can be represented as \( E(x) = E_v + e_s(x) \),

\[ E_v = E_{d\infty} \exp \left( -\frac{w_0 x}{\eta_0} \right) + \int_0^{+\infty} \frac{1}{2\pi i(\eta^2 - \eta_1^2)} \ast \left( C_0 + \frac{C_{-1}}{\eta - \eta_0} \right) \left( \frac{1}{X^+(\eta)} - \frac{1}{X^-(\eta)} \right) \exp \left( \frac{w_0 x}{\eta} \right) d\eta, \]

where

\[ E_{d\infty} = C_0 = \frac{\Lambda_1}{\Lambda_{\infty}}, \quad E_d = \frac{C_0(\eta_1/\eta_0^2 - \eta_1^2) + \alpha^-}{X(\eta_0)(\eta_1^{\alpha^+} - \eta_0^{\alpha^-})}, \]

\[ C_{-1} = -\frac{C_0[\eta_1 + \alpha^- (\eta_0^2 - \eta_1^2)]}{(\eta_1^{\alpha^+} - \eta_0^{\alpha^-})}, \quad \alpha^+ = \frac{X(\eta_1) \pm X(-\eta_1)}{2}, \]

\[ X(z) = \frac{1}{z} \exp V(z), \quad V(z) = \frac{1}{\pi} \int_0^{\infty} \frac{\zeta(\tau) d\tau}{\tau - z} \]

\[ \zeta(\tau) = \frac{1}{2i} \ln G(\tau) - \pi. \]

The detailed description of the function \( G(\tau) \) is given in [5]–[6].

Let us consider the electron response in a metal layer to an external ac electric field. We will calculate the electric field energy absorbed in a cylindrical area with the base area \( S \) and thickness \( a \). The external ac electric field \( E_0 \exp(-i\omega t) \) is applied perpendicular to the layer surface.

Absorption in a cylindrical volume with the base area \( S \) and thickness \( a \) is given by a well-known expression [7]

\[ Q = \frac{S}{2} \operatorname{Re} \int_0^a j(x) E^*(x) dx. \]

Here, \( j(x) \) is the current density and the asterisk denotes a complex conjugate.

Since we are considering a one-dimensional problem, the equation for the electric field has the form \( \frac{dE}{dx} = 4\pi q \) where \( q \) is the charge density. All quantities are assumed to depend on time as \( \exp(-i\omega t) \), i.e., \( E = E(x) \exp(-i\omega t) \). The continuity equation for the one-dimensional charge–current system is given by \( \frac{dJ}{dx} = -i\omega q(x) \).

Using some calculations, we obtain

\[ Q_s = \frac{\nu r S E_0^2}{8\pi \varepsilon} \operatorname{Im} \left\{ \frac{1}{w_0} \left[ C_0 + \frac{C_0(X(\eta_1) + X(-\eta_1))}{2X(\eta_1)X(-\eta_1)} \right] + \frac{2\eta_1}{\eta_1[X(\eta_1) - X(-\eta_1)][\eta_0^2 - \eta_1^2]} \right\} \]

\[ \times \left[ C_0 + \frac{\eta_0}{2}[X(\eta_1) + X(-\eta_1) - \eta_0][X(\eta_1) - X(-\eta_1)] \right] \]

The quantity \( Q_s \) corresponds to surface absorption. For a sufficiently broad plasma layer (with a width exceeding the electron mean free path), \( Q_s \) independent of the layer thickness.

Figure 1 shows the plots of surface absorption at \( \varepsilon = 0.0001 \). The curves 1, 2, 3 correspond to the values of the dimensionless chemical potential \( \alpha = -1, 0, 1 \). Figure 1 shows that as the chemical potential of the growth surface absorption increases.

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