Semiclassical theory of the circular photogalvanic effect in gyrotropic systems

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We develop a theory of circular photogalvanic effect (CPGE) for classically high photon energies which exceed the electron scattering rate but are small compared to the average electron kinetic energy. In this frequency range one can calculate the CPGE by using two different approaches. In the fully quantum-mechanical approach we find the photocurrent density by applying Fermi’s golden rule for indirect intraband optical transitions with virtual intermediate states both in the conduction and valence bands. In the framework of the semiclassical approach, we apply a generalized Boltzmann equation with accounts for the Berry-curvature induced anomalous velocity, side jumps and skew scattering. The calculation is carried out for a wurtzite symmetry crystal. Both methods yield the same results for the CPGE current demonstrating consistency between the two approaches and applicability of the semiclassical theory for the description of nonlinear high-frequency transport.

I. INTRODUCTION

The key signature of the Circular PhotoGalvanic Effect (CPGE) is the appearance of a photocurrent under illumination with circularly polarized light and reversal of its direction upon inversion of the light helicity. In time-reversal invariant systems the circular photocurrent is nonzero for point groups that allow optical activity or gyrotnropy. Among 21 crystal classes lacking inversion symmetry, only three noncentrosymmetric classes $T_d$, $C_{3h}$ and $D_{3h}$ are nongyrotropic. The CPGE was predicted by Ivchenko and Pikus [1] and Belinicher [2]. It was first observed and studied in Tellurium bulk crystals by Asnin et al. [3], see more references in the books [4–7]. Sometimes the term, “circular photocurrent”, is also referred to as “injection current” [8]. Renewed interest in the CPGE has been stimulated by prediction of its quantization in terms of the fundamental constants $\varepsilon$, $\hbar$ and the topological charge of Weyl nodes in the Weyl semimetals [9], see also Refs. [10–15].

The initial microscopic theories of the CPGE were based on the quantum-mechanical calculations applicable when the light frequency $\omega$ is much larger than the electron relaxation rate $\tau^{-1}$. The goal of the work [16] was to pave the way for the generalized semiclassical description of the circular and linear photogalvanic effects in solids, under conditions where the electron kinetic energy $\varepsilon$ exceeds the photon energy $\hbar\omega$. In this regime one can use the Boltzmann kinetic equation [17–20] with the field term containing the electric field of the electromagnetic wave.

Transport analysis of photocurrents in Ref. [16] showed that, within the semiclassical approach, the following mechanisms can contribute to the photogalvanic effects: (i) skew scattering which appears when going beyond the Born approximation [21–23], (ii) the electron anomalous velocity due to the Berry curvature in the electron Bloch wavefunction [17, 24, 25], (iii) the side-jump contribution to the electron velocity, and (iv) a change of electron energy caused by the side-jump in a scattering process in the presence of an external electric field [18, 23, 26, 27]. Since the mechanism (iii) is insensitive to the light helicity, only the other three mechanisms are relevant to the CPGE. Soon Moore and Orenstein [28] computed the Berry-phase contribution (ii) to helicity-dependent photocurrents in realistic circumstances, namely, for semiconductor quantum wells. This mechanism was then studied comprehensively for different materials and electron band models [29–41]. The Berry-curvature-related circular photocurrent has been expressed in an elegant form via the newly coined “Berry curvature dipole” [29].

In contrast to the Berry curvature-related current (ii), the mechanism (iv) has not attracted as much attention although it can lead to circular photocurrents of the same order of magnitude. Moreover, a similar mechanism has been studied in linear and nonlinear transport, the so-called anomalous-distribution mechanism of the anomalous Hall effect [23, 27, 42–44].

In this work, we present, within the same electron band structure model, both quantum-mechanical and semiclassical calculations of the intraband CPGE for a three-dimensional electron gas in the frequency range

$$\frac{\hbar}{\tau} \ll \hbar \omega \ll \varepsilon. \quad (1)$$

We will show that in this frequency range both approaches yield the same result which indicates consistency of the semiclassical approach. We will also discuss relative roles of above-mentioned contributions to circular photocurrents.

II. BAND STRUCTURE MODEL

We use the spin-independent band-structure model of a wurtzite-type semiconductor which includes the conduction band $\Gamma_{1c}$, and the valence bands $\Gamma_{6v}, \Gamma_{1v}$. The (real) Bloch functions at the $\Gamma$ point are labeled as $S (\Gamma_{1c}), X, Y (\Gamma_{6v})$ and $Z (\Gamma_{1v})$. In this basis the Kane
The parabolic dispersion in the conduction band is isotropic: $m_{c\perp} = m_{c||} \equiv m$ and 

$$\varepsilon_{c,k} = \frac{\hbar^2 k^2}{2m}.$$  

(7)

In order to simplify the computation of the photocurrent we further assume that the relation (6) is valid and use the isotropic electron dispersion in the conduction band. The Bloch functions at $k \neq 0$ are linear combinations of the basis functions $S, X, Y, Z$ multiplied by the exponential factor $e^{ikr}$, namely, 

$$\psi_{n,k}(r) = e^{ikr}u_{nk}, \quad u_{nk} = C_s S + C_x X + C_y Y + C_z Z.$$  

The coefficients $C_s, C_x, C_y$ and $C_z$ are usefully presented as a four-component column $C_{nk}$ which is an eigenvector of the matrix $H(k)$.

In the following we will calculate the Berry curvature $\Omega_{nk}$ and the elementary shifts $r_{nk,k'}$ of an electron in the $r$-space under the scattering between the states $(c,k)$ and $(c,k')$. For this purpose we expand the Bloch periodic amplitudes $u_{nk}$ in powers of $k$ up to the second order. For the conduction band, the expansion has the form 

$$u_{c,k} = C^{(c)} S + \left( -\frac{P_{\perp}}{E_g} - \frac{P_{\perp} Q k_z}{E_g E'_g} \right) (k_x X + k_y Y)$$  

$$+ \left( -\frac{P_{\parallel} k_x}{E_g} + \frac{P_{\parallel} Q k_x^2}{E_g E'_g} \right) Z ,$$  

where $E'_g = E_g + \Delta_e$ and the normalization coefficient equals 

$$C^{(c)} = 1 - \frac{1}{2} \left( \frac{P_{\perp}^2 k_x^2}{E_g^2} + \frac{P_{\parallel}^2 k_x^2}{E_g^2} \right).$$

The Bloch amplitude $u_{v_1,k}$ is a simple linear combination of $X$ and $Y$: 

$$u_{v_1,k} = \frac{k_y}{k_{\perp}} X - \frac{k_x}{k_{\perp}} Y.$$  

(9)

Similarly to $u_{c,k}$, the amplitudes $u_{v_2,k}$ and $u_{v_3,k}$ are linear combinations of three Bloch functions $S, (k_x X + k_y Y)/k_{\perp}$ and $Z$: 

$$u_{v_2,k} = \left( -\frac{P_{\perp} k_{\perp}}{E_g} + \frac{P_{\perp} Q k_{\perp} k_z}{E_g E'_g} \right) S$$  

$$+ C^{(v_2)} \left( \frac{k_x}{k_{\perp}} X + \frac{k_y}{k_{\perp}} Y \right) + \left( \frac{Q k_{\perp}}{E_g E'_g} - \frac{P_{\perp} P_{\parallel} k_z}{E_g E'_g} \right) Z ,$$  

$$u_{v_3,k} = \left( -\frac{P_{\parallel} k_x}{E'_g} + \frac{P_{\parallel} Q k_x^2}{E'_g E'_g} \right) S$$  

$$+ \left( \frac{Q k_z}{E'_g} + \frac{P_{\parallel} P_{\parallel} k_z}{E'_g E'_g} \right) (k_x X + k_y Y) + C^{(z)} Z .$$  

(11)

III. QUANTUM-MECHANICAL CALCULATION OF THE CIRCULAR PHOTOCURRENT

In the quantum-mechanical approach the intraband circular photocurrent is generated under indirect optical
transitions \((ck) \rightarrow (ck')\), with \(\varepsilon_{ck'} = \varepsilon_{ck} + \hbar \omega\), involving scattering by static defects and/or acoustic phonons. Here we focus on the elastic scattering processes and introduce the scattering matrix elements \(U_{n',nk}\), both intraband and interband. Then the compound matrix element of the indirect light absorption has the form

\[
M_{ck',ck} = \frac{V_{c,c}(k')U_{c,c}(k)}{\varepsilon_{ck'} - \varepsilon_{ck} + \hbar \omega} + \sum_{v=v_1,v_2,\ldots} \left( \frac{V_{v,c}(k')U_{v,c}(k)}{\varepsilon_{ck'} - \varepsilon_{ck} + \hbar \omega} \right),
\]

where \(V_{\nu',\nu}(k)\) is the optical matrix element of the direct transition \((nk) \rightarrow (n'k)\).

The first two terms in the rhs of Eq. (12) describe the indirect transitions via intermediate states in the conduction band \(c\) (intraband contribution) while the sum over \(v\) accounts for the virtual intermediate states in the valence band (interband contribution). The electric field of the incident light is defined as

\[
E(t) = E e^{-i \omega t} + E^* e^{i \omega t} = E_0 \left( e^{-i \omega t} e + e^{i \omega t} e^* \right).
\]

Then the intraband contribution to (12) can be written as

\[
M_{\text{intra}}^{ck',ck} = \frac{i e E_0}{m \omega} \frac{U_{c,c}(k')}{\varepsilon_{ck'} - \varepsilon_{ck} + \hbar \omega} [e \cdot (k' - k)],
\]

where \(e\) is the electron charge and the effective mass \(m\) is introduced in Eq. (7).

Within the accuracy of our calculation the electron energy dispersion and the photon energy in the denominators of the interband contribution can be neglected, and this contribution is reduced to

\[
M_{\text{inter}}^{ck',ck} = -\sum_{v} \frac{V_{v,c}(k')U_{v,c}(k)}{E_v - E_0},
\]

where \(E_0\) is the energy at \(k = 0\).

We assume that electrons are elastically scattered by the short-range defects. In this case the scattering matrix elements are given in the Born approximation by

\[
U_{n',nk} = U_0 \langle n'k | n_k \rangle,
\]

where \(U_0\) is real. Note that the perturbative expansion of the interband matrix element \(U_{ck',ck}\) or \(U_{ck',ck}\) starts from the first order in \(k\).

The optical matrix element is given by

\[
V_{\nu',\nu}(k) = \frac{ie E_0}{\omega} e \cdot v_{\nu',\nu}(k),
\]

where the velocity matrix elements are calculated according to

\[
v_{\nu',\nu}(k) = C_{\nu',k}^\dagger \hat{C}_{\nu,k} , \quad \hat{v} = \frac{1}{\hbar} \frac{\partial \hat{H}(k)}{\partial k}.
\]

Omitting tedious intermediate transformations we arrive at the following equation for the indirect optical matrix element

\[
M_{ck',ck} = i a e \cdot (k' - k) - b \left[ \varepsilon_z (k_{1z}' - k_{1z}) - \varepsilon_z (k_{1z} + k_{1z}) (k_{2z}' - k_{2z}) \right],
\]

where

\[
a = \frac{e E_0}{m \omega} U_0 , \quad b = \frac{e E_0 A}{2 \hbar \omega} U_0 , \quad A = \frac{2 P_l P Q}{E_g E_g} \left( \frac{1}{E_g} + 2 \right).
\]

The circular photocurrent is calculated according to [5]

\[
j^\text{CPGE} = 2 e n_i \frac{2\pi}{\hbar} \sum_{k'k} |v_{c,k'} \tau(\varepsilon_{k'}) - v_{c,k} \tau(\varepsilon_k)| \delta(\varepsilon_{k'} - \varepsilon_k - \hbar \omega),
\]

where the factor of 2 accounts for the double spin degeneracy and for brevity we set the sample volume to unity. The other notations are defined as follows: \(n_i\) is the defect concentration, \(v_{c,k} = \hbar k/m\) is the electron velocity in the conduction band, \(\tau(\varepsilon_k)\) is the energy-dependent momentum scattering time, \(f^0(\varepsilon_k)\) is the equilibrium electron distribution function, and we retain only the antisymmetric part of the squared matrix element

\[
|M_{ck',ck}|^2 = ab \left\{ (k_{1z}' - k_{1z}) [(k_{1z}' + k_{1z}) \times \kappa_z] + (k_{2z}' - k_{2z}) [(k_{2z}' + k_{2z}) \times \kappa_z] + 2 (k_{1z}' - k_{1z}) (k' \cdot k) \kappa_z \right\}.
\]

The last term in the rhs of Eq. (23) is proportional to \(\kappa_z\) and does not contribute to the current in agreement with the symmetry consideration (3). In the model under study the momentum relaxation rate has the form

\[
\frac{1}{\tau(\varepsilon_k)} = \frac{\pi}{\hbar} n_i U_0^2 g(\varepsilon_k) = \frac{\sqrt{2} m^{3/2}}{\hbar^4} U_0^2 n_i \sqrt{\varepsilon_k},
\]

where

\[
g(\varepsilon_k) = 2 \sum_{k'} \delta(\varepsilon_{k'} - \varepsilon_k)
\]

is the density of states. Thus, the exponent of the power-law dependence \(\tau(\varepsilon_k) \propto \varepsilon_k^n\) has the value

\[
\nu = \frac{d \ln \tau}{d \ln \varepsilon_k} = \frac{1}{2},
\]

In this paper we keep the general notation \(\nu\) to show the origin of the particular contributions to the photocurrent due to the derivative \(d\tau/d\varepsilon_k\). Only at the final stage we substitute the value \(\nu = -1/2\) into the equation for the photocurrent.

While performing the summation in Eq. (22) we take into account the energy conservation law \(\varepsilon_{k'} - \varepsilon_k = \hbar \omega\)
and use the Taylor’s linear expansion
\[ f^0(\varepsilon_{k'}) = f^0(\varepsilon_k) + \frac{df^0(\varepsilon_k)}{d\varepsilon_k} \hbar \omega, \]
\[ \tau(\varepsilon_{k'}) = \tau(\varepsilon_k) + \frac{d\tau(\varepsilon_k)}{d\varepsilon_k} \hbar \omega, \]
and the identities
\[ \frac{2\pi}{\hbar} \sum_k \delta(\varepsilon_{k'} - \varepsilon_k) = \frac{1}{n_i U_0^2} \tau, \]
\[ -2 \sum_k \varepsilon_k \frac{df^0}{d\varepsilon_k} = \frac{2}{N}, \]
where \( N = 2 \sum_k f^0(\varepsilon_k) \) is the electron density. The final result for the CPGE parameter \( \gamma \) reads
\[ \gamma = 2 \left( 1 + \nu \right) \frac{e^3 \hbar N}{\omega^2 \hbar}. \]

Now we turn to the semiclassical approach and show that the semiclassical circular photocurrent is also given by Eq. (28) with \( \nu = -1/2 \).

IV. SEMICLASSICAL DESCRIPTION OF CPGE

In this section we consider semiclassical contributions to \( \gamma \) mentioned in the Introduction and compare their sum with Eq. (28).

A. Berry curvature dipole (BCD) contribution

This contribution to the photocurrent is given by [16]
\[ j^{BCD} = \frac{2e^2}{\hbar} \sum_k [\Omega_{c,k} \times E(t)] f^{(1)}_{c,k}(t), \]
where we use the following notation. The Berry curvature \( \Omega_{c,k} \) is the curl of the Berry connection \( A_{c,k} \) (also called the Berry vector potential),
\[ \Omega_{c,k} = \frac{\partial}{\partial k} \times A_{c,k}, \quad A_{c,k} = i \left\langle u_{c,k} \left| \frac{\partial u_{c,k}}{\partial k} \right. \right\rangle, \]
and \( f^{(1)}_{c,k}(t) \) is the correction to the electron distribution function linear in the electric field \( E(t) \),
\[ f^{(1)}_{c,k}(t) = e^{-i\omega t} f^{(1)}_{c,k} + e^{i\omega t} f^{(1)*}_{c,k}. \]
In the framework of the classical Boltzmann equation in the relaxation-time approximation, it has the form
\[ f^{(1)}_{c,k} = -e\tau_\omega(\varepsilon_k) E_0 (e \cdot v_{c,k}) \frac{d\rho^0(\varepsilon_k)}{d\varepsilon_k}, \quad \tau_\omega = \frac{\tau}{1 - i\omega \tau}. \]

In the limit \( \omega \tau \gg 1 \), see Eq. (1), the “complex time” \( \tau_\omega \) is replaced by \( i/\omega \).

One can show using Eq. (8) for the periodic amplitude \( u_{c,k} \) that the Berry curvature is given, up to the first order, by
\[ \Omega_{c,k} = A(k \times c), \]
where \( c \) is the unit vector in the \( z \) direction and the coefficient \( A \) is defined by Eq. (21). An important point to stress is that the Berry curvature is proportional to the same parameter \( A \) which is present in Eq. (28) for the quantum-mechanical photocurrent. Substituting (13), (32) and (33) into Eq. (29) and summing over \( k \) we obtain
\[ \gamma^{BCD} = e^3 AN \frac{1}{\hbar^2 \omega}. \]

Comparing Eqs. (28) and (34) we see that the BCD contribution represents only a part of the total CPGE current and we need to consider other mechanisms of the semiclassical circular photocurrent.

B. Circular photocurrent due to the side-jump correction in the energy-conservation law

The kinetic equation for the electron distribution function \( f_{c,k} \) with allowance for the side-jumps is written as [16]
\[ \frac{\partial f_{c,k}(t)}{\partial t} + \frac{eE(t)}{\hbar} \frac{\partial f_{c,k}(t)}{\partial k} = n_i \sum_{k'} W_{k'k}(f_{c,k'} - f_{c,k}), \]
where
\[ W_{k'k}^{(0)} = \frac{2\pi}{\hbar} U_0^2 \delta(\varepsilon_{k'} - \varepsilon_k - eE(t) \cdot r_{k'k}), \]
\[ W_{k'k}^{(1)} = \frac{2\pi}{\hbar} U_0^2 \delta(\varepsilon_{k'} - \varepsilon_k), \]
and \( r_{k'k} \) is an elementary shift of the electron in the real space under the transition from \( (c,k) \) to \( (c,k') \). Equation (35) differs from the standard Boltzmann equation by the coordinate-shift correction to the collision integral. This correction follows from the change of the potential energy by \( -e(r_{k'k} \cdot E) \) under the scattering in the electric field due to the side-jump in which case the energy-conservation law reads [23, 42]
\[ \varepsilon_{k'} - \varepsilon_k - e(r_{k'k} \cdot E) = 0. \]

The side-jump, or the shift in real space due to scattering, is connected with the Berry vector potential by [23, 45, 46]
\[ r_{k'k} = -\frac{\text{Im} \left[ U_{c,k}^* (\nabla_k' + \nabla_k) U_{c,k'} \right]}{|U_{c,k'}|^2} + A_{c,k'} - A_{c,k}. \]
The matrix element of the elastic scattering \( U_{\mathbf{r}_k' \mathbf{k}} \) is calculated according to Eqs. (8) and (16) with the result

\[
\mathbf{r}_{\mathbf{k}/\mathbf{k}} = \frac{A}{2} \left[ (k_z - k'_z)(k'_\perp + k_\perp) + (k'_z - k_z) \mathbf{e} \right] \tag{39}
\]
determined by the same parameter \( A \) as the currents in Eqs. (28) and (34).

The photocurrent induced by the side-jump contribution to the collision integral (SJCI) is calculated according to

\[
\mathbf{j}^{\text{SJCI}} = 2e \sum_k \frac{\hbar}{m} g_k , \tag{40}
\]

where \( g_k \) is the steady-state correction to the electron distribution function proportional to \( E_0^2 \). It consists of two contributions \( g_k^0 \) and \( g_k^\alpha \).

1. **Field effect followed by SJCI**

The contribution \( g_k^0 \) is obtained by substituting the correction \( f^{(1)}_{\mathbf{k}'}(t) \) induced by the electric field, see Eqs. (31) and (32), into the rhs of Eq. (35) and averaging the product

\[
W_{\mathbf{k}'/\mathbf{k}}^{(1)}[f^{(1)}_{\mathbf{k}'}(t) - f^{(1)}_{\mathbf{k}}(t)]
\]

over time. This leads to

\[
g_k^0 = \tau n_\mathbf{k} \sum_{\mathbf{k}'} W_{\mathbf{k}'/\mathbf{k}}^{(1)}[f^{(1)}_{\mathbf{k}'}(t) - f^{(1)}_{\mathbf{k}}(t)] = -e\tau n_\mathbf{k} U_0^2 \tag{41}
\]

\[
\times \frac{2\pi}{\hbar} \sum_{\mathbf{k}'} |\mathbf{E}(t) \cdot \mathbf{r}_{\mathbf{k}'/\mathbf{k}}| \left[ f^{(1)}_{\mathbf{k}'}(t) - f^{(1)}_{\mathbf{k}}(t) \right] \frac{\partial \delta(\varepsilon_k - \varepsilon_{k'})}{\partial \varepsilon_{k'}},
\]

where the overline denotes a time average. We substitute \( f^{(1)}_{\mathbf{k}'}(t) \) from (31), (32) and \( \mathbf{r}_{\mathbf{k}'/\mathbf{k}} \) from (39) into Eq. (41), average over the time and the direction of \( \mathbf{k}' \) and then substitute \( g_k^0 \) into Eq. (40), average over the direction of \( \mathbf{k} \) and apply the identity (26) and the sum rule

\[
\sum_k F(\varepsilon_k) \frac{\partial \delta(\varepsilon_k - \varepsilon_{k'})}{\partial \varepsilon_{k'}} = -\sum_k \delta(\varepsilon_k - \varepsilon_{k'}) \frac{\partial}{\partial \varepsilon_k} \left[ \sqrt{\varepsilon_k} F(\varepsilon_k) \right], \tag{42}
\]

which is derived by integrating by parts and valid for an arbitrary differentiable function \( F(\varepsilon_k) \). This procedure allows us to reduce the contribution to the CPGE parameter \( \gamma \) to the sum

\[
\gamma_{\text{SJCl}}^a = -\frac{4e^3 A}{9\hbar^2 \omega} \sum_k \left[ 2(3 + \nu) \varepsilon_k \frac{d^2 f^{(2)}(\varepsilon_k)}{d \varepsilon_k^2} + \varepsilon_k^2 \frac{d^2 f^{(2)}(\varepsilon_k)}{d \varepsilon_k^2} \right].
\]

Taking into account Eq. (27) and the similar equation

\[
2 \sum_k \varepsilon_k^2 \frac{d^2 f^{(2)}(\varepsilon_k)}{d \varepsilon_k^2} = \frac{15}{4} N
\]

we arrive at

\[
\gamma_{\text{SJCl}}^a = e^3 A N (7 + 4\nu), \tag{44}
\]

2. **SJCI followed by the field effect**

The correction \( g_k^\alpha \) is obtained by first solving the kinetic equation without the field term but with the inhomogeneous term

\[
n_i \sum_{\mathbf{k}'} W_{\mathbf{k}'/\mathbf{k}}^{(1)} \left[ f^{(1)}_{\mathbf{k}'}(\varepsilon_{k'}) - f^{(1)}_{\mathbf{k}}(\varepsilon_k) \right]
\]

proportional to \( E_0 \). In the limit \( \omega \tau \gg 1 \), the solution reads

\[
f^{(1)}_{\mathbf{k}'}(t) = \frac{n_i}{\omega} \sum_{\mathbf{k}'} W_{\mathbf{k}'/\mathbf{k}}^{(1)} \left[ f^{(1)}_{\mathbf{k}'}(\varepsilon_k) - f^{(1)}_{\mathbf{k}}(\varepsilon_k) \right]. \tag{45}
\]

Secondly, the solution \( \tilde{f}^{(1)}_{\mathbf{k}'} \) is substituted into the field term of the kinetic equation. As a result we have

\[
g_k^\alpha = -\frac{e}{\hbar} \frac{\mathbf{E}(t) \cdot \nabla f^{(1)}_{\mathbf{k}'}(t)}{\varepsilon_k} \tag{46}
\]

Next we substitute (46) into Eq. (40) and integrate by parts to get

\[
\mathbf{j}^{\text{SJClb}} = \frac{2e^2}{m} \sum_k \left[ \mathbf{E}(t) \tilde{f}^{(1)}_{\mathbf{k}'}(t) \right] \frac{\partial [k_\alpha \tau(\varepsilon_k)]}{\partial \varepsilon_k}. \tag{47}
\]

The function in the sum averaged over time and the direction of \( \mathbf{k} \) can be reduced to

\[
(e \times \mathbf{E}) E_0^2 \frac{2e\nu e m A \partial f^{(0)}(\varepsilon_k)}{9\hbar^2 \omega} \varepsilon_k,
\]

which finally leads to

\[
\gamma_{\text{SJClb}} = \nu \frac{e^3 AN}{3h^2 \omega}. \tag{48}
\]

The sum of contributions to \( \gamma \) from the Berry curvature dipole, Eq. (34), and the side jumps in the collision integral, Eqs. (44) and (48), is given by

\[
\gamma_{\text{CPGE}} = \left( 1 + \frac{7 + 4\nu}{6} + \frac{\nu}{3} \right) \frac{e^3 AN}{h^2 \omega} = \left( \frac{13}{6} + \nu \right) \frac{e^3 AN}{h^2 \omega}.
\]

In the model under consideration this equation is valid only for \( \nu = -1/2 \) and reduces to

\[
\gamma_{\text{CPGE}} = \frac{5}{3} \frac{e^3 AN}{h^2 \omega}. \tag{49}
\]

The same value of \( \gamma \) follows from the quantum-mechanical result, Eq. (28), at \( \nu = -1/2 \).

V. **THE CIRCULAR PHOTOCURRENT DUE TO THE SKEW SCATTERING**

The skew-scattering mechanism is related to the antisymmetric part of the Boltzmann equation collision term arising beyond the Born approximation. Neglecting here
the side-jump term $W^{(1)}_{k'k}$ we can write the probability rate in noncentrosymmetric solids as

$$W^{(1)}_{k'k} = W^{(0)}_{k'k} + W^{(a)}_{k'k},$$

where the main term is defined in Eq. (36) and the antisymmetric term possesses the properties

$$W^{(a)}_{k'k} = -W^{(a)}_{-k',k} = -W^{(a)}_{kk'}.$$  

With allowance for this term the rhs of the kinetic equation (35) should be rewritten as

$$I\{f\} = n_i \sum_{k'} (W_{kk'} f_{ckv} - W_{k'k} f_{c'k}).$$

Its antisymmetric part can be simplified to

$$I_a \{f\} = n_i \sum_{k} W_{kk'}^{(a)} (f_{ckv} + f_{c'k})$$

$$= n_i \sum_{k'} W_{kk'}^{(a)} f_{c'k'},$$

where the sum property $\sum_{k'} W_{kk'}^{(a)} = 0$ is taken into account.

To the third order in $U$ the antisymmetric part of the elastic scattering probability is given by [4, 18, 47, 48]

$$W_{kk'}^{(a)} = \frac{(2\pi)^2}{h} \sum_{k''} U_{ckv} U_{c'k''} U_{ckv} U_{c'k''} U_{ckv} U_{c'k''},$$

$$= \frac{(2\pi)^2}{h} \delta(\varepsilon_{ckv} - \varepsilon_{c'k'})\delta(\varepsilon_{ckv} - \varepsilon_{c'k'}) .$$

In the band model (2) we find the following expression

$$W_{kk'}^{(a)} = \pi W_{kk'}^{(0)} g(\varepsilon_k) \frac{P_{ckv} P_{ckv'}}{E_g E_{g'}},$$

$$\times \left[ \left( \frac{k'^2}{E_{g'}} - \frac{2 k^2}{3 E_g} \right) (k' - k) + \frac{k'^2 k'^2}{E_{g'}} \right].$$

The PGE current is given by Eq. (40) where $g_k$ is replaced by the quadratic-in-$E_0$ correction, $f_{ck}^{(2a)}$, averaged over time. It is calculated in three steps. First, we find the first-order correction $f_{ck}^{(1a)}$, see Eq. (31). Then we substitute it into the collision integral and determine a correction, $f_{ck}^{(1a)}$, to the distribution function induced by the antisymmetric part (50)

$$f_{ck}^{(1a)} = \tau_\omega I_a \{f^{(1)}\} .$$

Lastly, we find the desired second-order correction

$$f_{ck}^{(2a)} = -\frac{eE}{\hbar} \frac{\partial f_{ck}^{(1a)}}{\partial k} .$$

The photocurrent can be rewritten as

$$j = 2e \sum_k \nu_{ck} f_{ck}^{(2a)} = \frac{2e^3 E_0^2}{h} \sum_k \nu_{ck} \tau$$

$$\times \left( e \cdot \frac{\partial}{\partial k} \right) \left[ \tau^2 I_a \left\{ e^* \cdot \nu_{c'k} \frac{\partial f_0(\varepsilon_k)}{\partial \varepsilon_{c'k}} \right\} \right] + c.c.$$  

In the limit $\omega \tau \rightarrow \infty$ when $\tau^2 \approx -\frac{1}{\omega^2} (1 + \frac{21}{\omega^2})$ the circular photocurrent vanishes and one needs to take into account the next-order correction in the expansion

$$\tau^2 \approx -\frac{1}{\omega^2} \left( 1 + \frac{21}{\omega^2} \right) .$$

Omitting the details, we present the result for the degenerate electron gas

$$\gamma_{\text{skew}} = -\frac{8\pi e^2 A'}{15 \hbar^2 \omega} \varepsilon_F g(\varepsilon_F) U_0 \frac{1}{\omega \tau} ,$$

where $\varepsilon_F$ is the Fermi energy, $\tau = \tau(\varepsilon_F)$,

$$A' = \frac{P_{ckv}}{E_g E_{g'}} \left[ \frac{1}{E_g} + \frac{2}{3} \frac{E_{g'}}{E_{g'}} \right],$$

and the dimensionless parameter $g(\varepsilon_F) U_0$ is assumed to be small which is the criterion for validity of the expansion of the collision term in powers of $U$ leading to Eq. (51). Note that the ratio of $\gamma_{\text{skew}}$ to the $\gamma_{\text{CPGE}}$ value of Eq. (49) is a product of a large parameter $\varepsilon_F \hbar \omega$ and two small parameters, $g(\varepsilon_F) U_0$ and $A_{\text{CPGE}}^{-1}$. It is interesting to mention that the dependence $\gamma_{\text{skew}} \propto 1/\omega^3$ is typical for the skew-scattering induced CPGE at high frequencies [49, 50]. It should also be stressed that the CPGE photocurrent (55) arises due to the $(\omega \tau)^{-1}$ correction in Eq. (54). This means that the result (55) cannot be obtained in the quantum-mechanical approach used in Sect. III in the limit $\omega \tau \rightarrow \infty$, and one needs to apply the Green’s function diagram technique which is beyond of this study.

According to Eq. (3) the macroscopic parameter $\gamma$ relates the variables $j$ and $\mathbf{E} \cdot \mathbf{E_0}$ which both change the sign under the time reversal $T$. This explains why the expression $\gamma_{\text{CPGE}}$ in Eq. (49) does not contain dissipative parameters. The expression for $\gamma_{\text{skew}}$ agrees with the time inversion symmetry considerations because the presence of the dissipative factor $\tau^{-1}$ is compensated by an extra $\delta$-function in Eq. (51).

The anomalous Hall coefficient also allows for a fourth-order correction, $U_4^2$, to the antisymmetric part of the scattering rate [23]. An estimate for $\omega \tau \gg 1$ shows that this skew contribution to $\gamma$ is smaller than that in Eq. (49) by $(\omega \tau)^2$ and can be neglected.

VI. CONCLUDING REMARKS

The semiclassical approach in the nonlinear transport is based on the two conditions: (i) the photon energy $\hbar \omega$ is much smaller than the typical electron energy $\varepsilon$, and (ii) in the kinetic equation for the free carrier distribution function, the standard or generalized Boltzmann equation, the field term

$$\frac{\partial f_{ck}}{\partial k} \frac{\partial f_{ck}}{\partial k}$$  

(56)
contains the electric and magnetic fields of the light wave as follows

\[ \hbar \dot{c} = e \left( E + \frac{\dot{c}}{c} \times B \right), \]

where \( \dot{c} \) is the generalized semiclassical velocity \cite{17} and

\[ \dot{c} = \frac{\hbar}{m} + \frac{\hbar}{\Omega} \times E. \]

Both the previous and present study show that the challenge in calculating a second-order response to electromagnetic waves is related to the choice of the collision term. The latter has corrections due to the side-jump, Sect. IV.B, and skew, Sect. V, scatterings.

The alternative quantum-mechanical approach is to calculate the optical transition matrix elements linear in the electric or magnetic field of the light wave. Microscopically, the total photocurrent density is generally a sum of two different contributions, ballistic and shift currents, \( j = j_b + j_s \) \cite{51}. To calculate the ballistic photocurrent one needs to solve the Boltzmann equation where the electromagnetic field is present in the generation rate proportional to the squared transition matrix element, calculated with allowance for the asymmetry of the electron-photon and/or electron-phonon or electron-impurity interaction. The shift photocurrent is determined by a product of the light-induced quantum transition probability rate and the elementary displacement (side-jump) \( r_{mk'} nk \) of a free carrier at the moment of the transition.

In the present work we have aimed to reconcile the two approaches. It is possible to do in the frequency range (1). Firstly, the agreement between results (28) and (49) for the circular photocurrent confirms the validity of the theory. Secondly, we have shown that the agreement is achieved only if, in the semiclassical approach, one takes into account the Berry curvature dipole mechanism and the contribution from the field-induced side-jump correction to the transferred electron energy.

ACKNOWLEDGMENTS

The work of L.E.G. was supported by the Foundation for the Advancement of Theoretical Physics and Mathematics “BASIS”.

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