Effects of anisotropy and disorder on the conductivity of Weyl semimetals

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We study dc conductivity of a Weyl semimetal with uniaxial anisotropy (Fermi velocity ratio \(\xi = v_1/v_\parallel \neq 1\)) considering the scattering of charge carriers by a wide class of impurity potentials, both short- and long-range. We obtain the ratio of transverse and longitudinal (with respect to the anisotropy axis) conductivities as a function of both \(\xi\) and temperature. We find that the transverse and longitudinal conductivities exhibit different temperature dependence in the case of short-range disorder. For general long-range disorder, the temperature dependence (\(\sim T^4 \)) of the conductivity turns out to be insensitive of the anisotropy in the limits of strong (\(\xi \gg 1\)) and weak (\(\xi \approx 1\)) anisotropy.

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I. INTRODUCTION

Weyl semimetals (WSM) are three-dimensional (3D) analogs of graphene.\(^1\) Their quasiparticles are described by the massless 3D Dirac Hamiltonian. Such systems were first proposed as an exotic theoretical possibility, with an expectation to observe some of their features in pyrochlore iridates\(^3\) and in certain semiconductor heterostructures.\(^4\) Recent experiments uncovered several chemical compounds, which can be classified as WSM\(^5\). The subject is attracting growing theoretical and experimental interest. (see, e.g., the reviews in Refs. 1\textsuperscript{2},14).

Charge transport in WSM has also received considerable attention and a number of interesting phenomena have been discovered (see Ref. 1\textsuperscript{14} and references therein). The main emphasis was on manifestations of topological effects: an additional topological protection of the gapless spectrum near the Dirac points in 3D and the chiral anomaly.\(^15\textsuperscript{2}–22\) as well as an unusual Kondo effect.\(^22\) Disorder and impurity effects have also attracted significant attention.\(^23\textsuperscript{2}–26\) A detailed study of the influence of Coulomb disorder in highly compensated WSM was undertaken in Ref. 27. The problem is that even in the simplest case of delta-correlated disorder, the field theory of a WSM is non-renormalizable. Progress in overcoming this stumbling block was recently achieved in Refs. 28\textsuperscript{2}–29\textsuperscript{2}, where the conductivity of weakly-disordered semimetals was treated in terms of the \(\epsilon\)-expansion within the renormalization group (RG) approach. However, Weyl semimetals correspond to \(\epsilon = -1\) and the predictions are qualitative. Nevertheless, it appeared to be possible to reveal some specific features of the conductivity for different kinds of disorder and even to predict a disorder-driven quantum phase transition distinct from the Anderson transition. A RG approach\(^30\) has been also applied to demonstrate the possibility of non-Fermi-liquid behavior in systems with isotropic 3D Dirac points having a weak disorder and Coulomb-type interparticle interactions.\(^30\)

The computation of the isotropic conductivity was recently performed\(^25,31,32\) in the framework of the kinetic-equation technique, improved by the self-consistent Born approximation, as well as numerically.\(^33\)

Most Weyl semimetals obtained in laboratories so far are anisotropic.\(^2\)\textsuperscript{28} The anisotropy can be also induced by a linearly-polarized electromagnetic wave.\(^24\) Some effects related to the anisotropy and tilting of the Dirac cones have recently been treated.\(^25\) However, the influence of anisotropy on the charge transport in WSM with quenched disorder has remained unexplored so far. In this paper, we analyze the effects of disorder and anisotropy on the conductivity, in the framework of the Born approximation. We treat these effects in the diagrammatic approach, both for short- and long-range disorder taking into account the uniaxial anisotropy. We obtain analytical results for the cases of strong and weak anisotropy. The diagrammatic framework allows us to find out the validity range of the Fermi-liquid approach in WSM.

The potential produced by impurities (disorder potential) is introduced as a rather general spatial distribution \(u(x)\). Its characteristic scale in momentum space is specified by a parameter \(p_0\). Thus, at small values of \(p_0\), we are dealing with long-range disorder, whereas large \(p_0\) corresponds to short-range disorder. The ratio of Fermi velocities perpendicular and parallel to the \(z\) axis, \(\xi = v_\perp/v_\parallel\), becomes an additional control parameter of the problem.

We have found that for sufficiently low temperatures (or doping level \(\epsilon_F\)), when the disorder can be considered to be a short-range one (\(p_0 \gg \max\{T/v_\parallel, \epsilon_F/v\}\)), the longitudinal and transverse conductivities exhibit a different temperature dependence

\[
\frac{\sigma_\parallel \xi^2}{\sigma_\perp} = 1 - \frac{4}{15} \frac{dg(0)}{\sqrt{x^2}} \left(1 + \xi^{-2}\right) \frac{\pi^2 T^2 / 3 + \epsilon_F^2}{(v \parallel p_0)^2},
\]

where \(g(p^2/p_0^2)\) is the impurity structure factor and \(dg(0) \equiv dg(x)/dx|_{x=0}\). The temperature dependence in
Eq. (1) is in fact the first term of the asymptotic series in \( \max\{T/\|v\|, \varepsilon_F/\varepsilon\}/p_0 \ll 1 \). This dependence becomes even more pronounced at higher temperatures \( T \sim p_0/|v| \).

The \( \sigma_{||}/\sigma_\perp \) ratio saturates to a temperature-independent constant in the high-temperature (or long-range disorder) limit, when \( (p_0 \ll \max\{T/|v|, \varepsilon_F/\varepsilon\}) \). At \( \varepsilon_F \ll T \), the temperature dependence of both components of the conductivity obeys the relation

\[
\sigma_{||}, \sigma_\perp \sim T^4
\]

regardless of the particular form of the disorder potential or impurity structure factor. In the long-range disorder limit, for \( \xi \gg 1 \), \( \xi \approx 1 \), and \( \xi \ll 1 \), we found that the \( \sigma_{||}/\sigma_\perp \) ratio approaches constant values, independent of temperature and disorder potential. The results concerning the conductivity ratio can be summarized in the form

\[
\frac{\sigma_{||}}{\sigma_\perp} = \begin{cases} 
\frac{3}{2} \left( \frac{1}{2} + 4 \ln 2 \right) + O(\xi^{-1}), & \xi \gg 1, \\
1 + O(\delta \xi^2), & \xi = 1 + \delta \xi, \\
c_3 + O(\delta \xi^2), & \xi \ll 1,
\end{cases}
\]

where \( c_3 \) is a constant of the order of unity.

The scales of the problem are the temperature \( T \) of the system, Fermi energy \( \varepsilon_F \), and the elastic scattering rate \( 1/\tau \). The latter is assumed to be small, so

\[
\frac{1}{\max\{\varepsilon_F, T\}/\tau} \ll 1
\]

is a small parameter of the problem. Our task is to compute the Drude conductivity. The fermion doubling theorem implies that the WSM spectrum always has an even number of Weyl points. For simplicity, we assume that the charge carriers have identical spectra near all Weyl points and compute the conductivity per point. All the results for the conductivity should be multiplied by the number of Weyl points.

The disorder potential is assumed to be quite weak, so that the Born approximation is justified

\[
|u(r)| \ll \min\{v_{||}, v_\perp\}/p_0.
\]

Note here that our results are applicable not only to the Weyl semimetals themselves, but to a wider class of materials with the 3D Dirac spectrum (topological Dirac semimetals etc., see the classification given in Ref. [36]).

In Section II, we describe the formalism used in dealing with the scattering problem, give all the necessary definitions, and describe the employed calculation procedures. In this section, we also analyze the limits of applicability of the Fermi-liquid approach in the case of WSM. In Section III, we revise the general temperature dependence of the conductivity in isotropic Weyl semimetals with long- and short-range disorder. Section IV deals with the anisotropic case. We provide a detailed analysis of the solution to the Dyson equation for the singular part of the vertex function for short- and long-range disorder potentials. We compute the conductivity for different \( T/\varepsilon_F \) ratios and different values of the anisotropy parameter \( \xi \). The conclusions are presented in Section V. Several important technical issues are presented in the Appendices.

II. FORMALISM

A. The model

We consider a system with a Hamiltonian, which generally has uniaxial anisotropy. Namely, the Fermi velocity along a specified axis \( n_0 \) (the longitudinal component) is different from that in the perpendicular direction \( v_\perp = \xi v_{||} \).

\[
H = H_0 + H_{\text{dis}}; \\
H_0 = -i \int \psi^\dagger(r) \left[ \sigma_{||} \partial_{r_{||}} + v_\perp \sigma_\perp \partial_{r_\perp} \right] \psi(r) dr,
\]

where \( u(r) \) is the disorder potential, \( \psi \) are quasiparticle field operators, and \( \sigma_{||} = \tau_n n_0, \psi_{||} = \tau_\parallel, \psi_\perp = \sigma - \sigma_n n_0, r_\parallel = r - n_0 (n_0 r) \) are the projections of Pauli matrices and radius vectors. The disorder potential correlation function reads

\[
\int dr e^{-i p r} \langle u(r) u(0) \rangle = \frac{n_{\text{imp}} u_0^2}{p_0^6} g \left( \frac{L^2}{p_0^2} \right),
\]

where \( g(p^2/p_0^2) \) is the dimensionless Fourier transform of the normalized \( (g(0) = 1) \) disorder structure factor. In our case, it incorporates the correlation of impurity positions, as well as the form of the potential. The parameter \( u_0 \) plays the role of the amplitude of the disorder potential and \( n_{\text{imp}} \) is the concentration of impurities. The potential is assumed to be isotropic. The last assumption certainly depends on the nature of the disorder. For example, the isotropic Coulomb potential may acquire an anisotropic screening by charged carriers with an anisotropic spectrum. Here, we neglect the induced anisotropy of the potential. Our task is to emphasize the main effect of the anisotropic spectrum on the transport properties, which manifests itself even with an isotropic potential.

To simplify the analysis, we rescale the coordinates and \( \psi \) operators to absorb the anisotropy in \( H_0 \) according to

\[
r_{||} = r_{||}', \quad r_\perp = \xi r_\perp', \quad \psi(r) = \frac{1}{\xi} \psi'(r').
\]

Then, the Hamiltonian changes its form to

\[
H_0' = -i v_\parallel \int \psi^\dagger(r') (\sigma_\parallel \partial_{r_\parallel}) \psi'(r') dr', \\
H_{\text{dis}}' = \int dr' \psi^\dagger(r') u(r_\parallel' + \xi r_\perp') \psi'(r'),
\]

where
which leads to the modified disorder correlation function
\[
\frac{1}{\hbar} \langle u(r'_e + \xi'_{\perp}) u(0) \rangle = \frac{n_{\text{imp}} q_0^2}{\hbar \xi^2} g' \left( \frac{p^2}{p_0^2} \right),
\]
where we have decomposed the momentum as: \( p^2 = p_0^2 + (p^2 - p_0^2) \rightarrow (p_0^0)^2 + \xi^{-2}[p^2 - (p_0^0)^2] \).

The Feynman rules are extracted from (9) and (10), and depicted in Fig. 1.

\( \varepsilon, q \) \( \rightarrow \) \( \frac{1}{\varepsilon - \nu \| p \sigma + i \nu} \)
\( q, -q \) \( \leftrightarrow \) \( \frac{n_{\text{imp}} q_0^2}{\hbar \xi^2} g \left( \frac{1 - \xi^{-2}}{p_0^2} \right) \)

The conductivity tensor is found via the Kubo formula
\[
\sigma_{\alpha \beta}(\omega, 0) = \frac{e^2 \Pi^R_{\alpha \beta}(\omega)}{i \omega},
\]
\[
\Pi_{\alpha \beta}(\omega) = i \int dt \, dr \langle [j_\alpha(t, r), j_\beta(0, 0)] \rangle e^{i \omega t} \theta(t),
\]
where \( \Pi^R_{\alpha \beta}(\omega) \) is the retarded polarization operator and \( j_\alpha(t, r) = \psi^\dagger(t, r) \sigma_\alpha v_\alpha \psi(t, r) \) is the quasiparticle current operator. The averaging \( \langle \ldots \rangle \) is assumed to be done over the Gibbs distribution as well as over different realizations of disorder. We will be interested in the system response to a uniform electric field, constant in time. For this purpose, we set \( q = 0 \) and \( \omega \rightarrow 0 \).

\( \Pi_{\alpha \beta}(\omega) = \frac{1}{2} \frac{J_\alpha}{\omega} \begin{array}{c} \varepsilon + \omega, p \\ \varepsilon, p \end{array} + \frac{1}{2} \frac{J_\beta}{\omega} \begin{array}{c} \varepsilon + \omega, \varepsilon, p \end{array} \)

\( G_\alpha(\varepsilon, q) = \begin{array}{c} \varepsilon, q \end{array} \quad \text{exact Green's function} \)

\( J_\alpha(\varepsilon, \varepsilon + \omega, \omega; p) = \begin{array}{c} \varepsilon, p \end{array} \quad \text{exact vertex} \)

The computation of the polarization operator in the lowest order of the \( 1/(\varepsilon_F \tau) \) expansion involves the summation of disorder ladder series (accounting for the difference between averages \( \langle j_\alpha J_\beta \rangle \) and \( \langle j_\alpha \rangle \langle j_\beta \rangle \)) as well as one-loop corrections to the Green’s functions (mainly responsible for the finite quasiparticle lifetime).

Due to the Onsager relations, the conductivity (as well as the polarization operator) is a symmetric tensor. Thus, the exact diagrammatic representation for \( \Pi_{\alpha \beta}(\omega) \) (we denoted \( \Pi_{\alpha \beta}(\varepsilon, 0) \equiv \Pi_{\alpha \beta}(\varepsilon) \)) allows for its symmetric form (see Fig. 2).

The non-interacting and disorder-averaged Green’s functions are related via the standard diagrammatic equation depicted in Fig. 3 and have the form
\[
G^R(\varepsilon, q) = \left[ \varepsilon - \nu \| p \sigma - \Pi^R(\varepsilon, q) \right]^{-1}.
\]

The self-energy is worked out in the Born approximation; its diagrammatic representation is shown in Fig. 3b.

The ladder summation in the polarization operator is included into the renormalized vertex \( J \) corresponding to the electric current. To perform the summation, one has to solve the corresponding Dyson equation depicted in Fig. 4.

The conductivity tensor in a system with the uniaxial anisotropy is characterized by only two eigenvalues \( \sigma_\parallel, \sigma_\perp \) and in “principal rescaled axes” takes the form
\[
\sigma_{\alpha \beta} = \begin{pmatrix} \varepsilon^2 \sigma_\parallel & 0 & 0 \\ 0 & \sigma_\perp & 0 \\ 0 & 0 & \sigma_\perp \end{pmatrix}.
\]

The general expression for the conductivity in the leading \( 1/(\varepsilon_F \tau) \) approximation reads
\[
\sigma_{\alpha \beta}(T) = \frac{1}{4T} \int \frac{d\varepsilon}{2\pi} \frac{\sigma_{\alpha \beta}(\varepsilon)}{\cosh^2 \frac{\varepsilon}{2T}}.
\]
\[ \sigma_{\alpha \beta}(\varepsilon) = \frac{e^2 v^2}{4\pi} \int \frac{dp}{(2\pi)^3} \text{tr} \left\{ G^A(\varepsilon, p) [J^{ARR}_\alpha(\varepsilon, p) - \sigma_\alpha] G^R(\varepsilon, p) \sigma_\beta + G^A(\varepsilon) [J^{ARR}_\beta(\varepsilon, p) - \sigma_\beta] G^R(\varepsilon, p) \sigma_\alpha \right. \\
\left. - [G^R(\varepsilon, p) - G^A(\varepsilon, p)] \sigma_\alpha [G^R(\varepsilon, p) - G^A(\varepsilon, p)] \sigma_\beta \right\}. \] (15)

Here, for brevity, we denoted \( J^{ARR}_\alpha(\varepsilon, 0; p) \equiv J^{ARR}_\alpha(\varepsilon, p) \), and \( J^{ARR} \) is the singular part of the vertex function (see Appendix A for details).

C. Validity of the Fermi-liquid approach

Expression (15) for \( \sigma_{\alpha \beta}(\varepsilon) \) is rather complicated and deserves special attention. The integrand defining \( \sigma_{\alpha \beta}(\varepsilon) \) is deliberately split into two lines. The second line leads to a convergent integral because \( (G^R - G^A) \sim 1/p^2 \), when \( p \to \infty \). The first line of (15) contains the vertex \( J^{ARR} \), which makes it problematic even in the isotropic case (\( v_\parallel = v_\perp = v \)). At first glance, the expression seems suitable enough, because the product \( G^R G^A \) is sharply peaked at \( p = p_c = \varepsilon / v \). The integrand therefore, appears to be completely determined by the vicinity of \( p = p_c \). However, a more attentive look reveals that the product \( G^R G^A \) converges rather slowly: \( G^R G^A \sim 1/p^2 \) at \( p \ll p_c \ll \tau \varepsilon p_c \), and \( G^R G^A \sim 1/p^4 \) at \( p > \tau \varepsilon p_c \). Thus, we need to know at least the asymptotic behavior of \( J^{ARR}(\varepsilon, p) \) at large momentum in order to complete the calculation.

This difficulty is closely related to the justification of the Fermi-liquid approach. Indeed, this justification includes two constraints: (a) the smallness of the quasiparticle scattering rate guarantees the applicability of perturbation theory (in most cases, the vacuum state, the Fermi sphere, survives); and (b) all observables are determined by the scattering processes in the proximity of the Fermi sphere. The latter statement allows expanding the momentum near the Fermi surface in every integration \( (dp = (\varepsilon_F / v)^2 dp \) in 3D) making any momentum integral one-dimensional and fast converging.

The justification of the second constraint follows from the fact that a momentum integral defining the observable always involves the \( G^R(p) G^A(p) \) term (sharply peaked near the Fermi sphere). However, we have just seen that the latter argument fails when computing the WSM conductivity. The integrand defining the conductivity does contain a sharp \( G^R(p) G^A(p) \) peak as well as a long tail (see Fig. 5).

To justify the Fermi-liquid framework, we need to accurately estimate the contribution given by the tail. We divide the integration domain according to \( p^2 dp = \)

\[ (p^2 - p_c^2) dp + p_c^2 dp. \] Then, \( F^{\text{sing}} \) is split into two parts

\[ F^{\text{sing}} = I + II, \]

\[ I = \int (p^2 - p_c^2) G^A(\varepsilon, p) [J^{ARR}_\alpha(\varepsilon, p) - \sigma_\alpha] G^R(\varepsilon, p) \sigma_\beta dp, \]

\[ II = p_c^2 \int G^A(\varepsilon, p) [J^{ARR}_\alpha(\varepsilon, p) - \sigma_\alpha] G^R(\varepsilon, p) \sigma_\beta dp. \] (16)

Part II is already convergent. The convergence of part I entirely depends on the asymptotic behavior of the vertex function. The ultraviolet behavior of the vertex is discussed in Appendix C, where it is proved to decay fast enough to secure its convergence. Part I is estimated in Appendix D, where it is shown that

\[ I \sim \max \left\{ 1, \frac{p_0}{p_c(1/\varepsilon)} \right\} \frac{1}{\varepsilon \tau} II. \] (17)

This means that the Fermi-liquid approach holds with 1/(\( \max[\varepsilon_F, T] \tau \)) accuracy. Therefore, some care needs to be taken when working out the weak-localization corrections (WLC) to conductivity in, e.g., the 3D case. Indeed, this correction is determined by the next term in the 1/(\( \varepsilon_F \tau \)) expansion, and the WLC in 3D reads \( \delta \sigma_{\text{WLC}} / \sigma \sim 1/(\varepsilon_F \tau) \). Therefore, when computing corrections to the conductivity in the 3D case, one has to take into account the values of the momentum far from the Fermi surface.

As a result, expanding all the momentum integrals near the Fermi surface, we can omit (in the leading order) the

![Figure 5: Contributions to the conductivity. Region I represents the Fermi liquid contribution; region II originates from the scattering far from the Fermi surface.](image-url)
$G^R G^R$ and $G^A G^A$ terms in the expression \[ \frac{\sigma_{\alpha\beta}(\epsilon)}{4\pi} = \frac{e^2 v^2}{4\pi} \int \frac{dp}{(2\pi)^3} \left\{ G^A(\epsilon, p) J^{ARR}_\alpha(\epsilon, p) G^R(\epsilon, p) \sigma_{\beta} + G^A(\epsilon) J^{ARR}_\beta(\epsilon, p) G^R(\epsilon, p) \sigma_{\alpha} \right\} . \] (18)

III. CONDUCTIVITY IN THE ISOTROPIC CASE

In order to understand how to tackle the anisotropic problem, we briefly outline the diagrammatic derivation of the isotropic conductivity. Wherever isotropy is implied, we omit the \( \parallel \) and \( \perp \) indexes, and write \( v_\parallel = v_\perp = v. \) In the corresponding limits, we reproduce the earlier results for the conductivity.\[ \text{(20)} \]

A. Disorder averaging

The self-energy of the Green’s function has matrix structure (due to the presence of \( \sigma \)-matrices) and consists of two parts: \( \Sigma = \Sigma^R + \Sigma^I. \) The first part is responsible for the renormalization of Fermi velocities and the rescaling of the Fermi field operators; the second part is responsible for the non-vanishing relaxation rate. Perturbation theory can be applied only if \( |\Sigma| \ll \max\{|\epsilon, T\}. \) The estimate gives

\[
\frac{|\Sigma|^R}{\epsilon} \sim \left( \frac{v_0}{\epsilon} \right) \frac{1}{\tau} \sim \lambda(\rho_0) \min \left\{ 1, \left( \frac{p_0 v}{\epsilon} \right)^2 \right\},
\]

\[
\frac{1}{\tau} \sim \left( \frac{n_{\text{imp}} u_0^2}{v^3 p_0^6} \right) \min \left\{ \frac{\epsilon^2}{v^3 p_0^6}, \left( \frac{p_0 v}{\epsilon} \right)^2 \right\},
\]

where \( |\Sigma|^R \) is understood as the modulus of any of its matrix component, \( \epsilon \sim \max\{|T/v, \epsilon_F/v\} \) (\( \epsilon_F \) can be considered here as the the doping level). The dimensionless coupling constant

\[
\lambda(\rho_0) = \frac{n_{\text{imp}} u_0^2}{v^3 p_0^6}
\]

represents the strength of the potential. It is defined at the characteristic disorder scale \( \rho_0. \)

Clearly, despite the smallness of the Fermi-liquid parameter \( 1/\epsilon\tau, \) the first part of the self energy may lead to a strong renormalization for short-range \( (\rho_0 \gg \epsilon/v) \) disorder, leading on the initial coupling strength \( \lambda(\rho_0). \) This is in contrast to the situation in ordinary metals, where the ultraviolet cut-off \( \rho_0/\epsilon_F \sim 1 \) and the smallness of the \( 1/\epsilon\tau \) parameter guarantees the validity of perturbation theory. In WSM, it is appropriate to consider both limits: short- \( (\rho_0 \gg \max\{p_F, T/v\}) \) and long-range \( (\rho_0 \ll \max\{p_F, T/v\}) \) disorder.

1. Renormalization due to short-range disorder

The analysis of the influence of strong short-range disorder \( \lambda(\rho_0) \ll 1 \) was attempted in Ref. 25 via the self-consistent Born approximation (SCBA). The latter however overlooks the diagrams with crossed disorder lines. In WSM, these diagrams happen to be of the same order as the diagrams included in the SCBA. Therefore, the SCBA is an uncontrolled approximation for a WSM. A fully consistent analysis was performed in Refs. 28-29 (by means of the \( \epsilon \)-expansion). Crucially, it was shown that the disorder operator becomes relevant if the initial coupling exceeds some critical value

\[
\lambda(\rho_0) > \lambda_* \sim 1.
\]

For \( \lambda(\rho_0) > \lambda_* \), the disorder coupling grows with decreasing momentum, which leads to the metal–insulator transition. If the initial disorder strength is small, \( \lambda(\rho_0) \ll \lambda_* \), then the disorder operator is irrelevant. The corresponding running value of the dimensionless coupling constant is

\[
\lambda(p) = \frac{\lambda(\rho_0) p}{p_0},
\]

where \( p \sim \max\{\epsilon_F/v, T/v\} \). We will focus on the case \( \lambda(\rho_0) \ll 1 \) and discard the renormalization of the Fermi velocity and field operators.

2. Long-range disorder

As one can deduce from Eq. \[ \text{(18)} \], the long-range \( (\rho_0 \ll \epsilon/v) \) disorder does not present a problem since the corresponding \( \Sigma_1 \) part is much smaller than the \( 1/(\epsilon\tau) \) parameter. Therefore, here we also discard the renormalization of the Fermi velocity and field operators.

B. The general Dyson equation and its solution in the isotropic case

Surprisingly, even for an anisotropic system, the Green’s function preserves its simple isotropic form (in the limiting cases of short-range and long-range disorder, see Appendix F with detailed calculations). The Green’s function has the form \[ \text{(23)} \]

\[
G^{R,A}(\epsilon, p) = \frac{\epsilon + v|p|\sigma}{(\epsilon + v|p|\mp \epsilon/F \tau)^2}.
\]
We will need the following scattering times
\[
\tau (\mathbf{m}_0) = \frac{n_{imp}^2}{2\tau_1^2} \int \frac{d\Omega}{4\pi} g_{p,(n-n')} (n,n_0),
\]
\[
\tau (\mathbf{m}_0) = \frac{n_{imp}^2}{2\tau_1^2} \int \frac{d\Omega}{4\pi} g_{p,(n-n')} (1 - \cos \theta_{nn'})
\]
\[
= \frac{1}{\tau - \tau_1},
\]
\[
\tau (\mathbf{m}_0) = \frac{n_{imp}^2}{2\tau_1^2} \int \frac{d\Omega}{4\pi} g_{p,(n-n')} (1 - \cos^2 \theta_{nn'})
\]
\[
= \frac{1}{\tau - \tau_2}.
\]
(24)
The derivation of the scattering rates is sketched in Appendix B. In accordance to what was said about the validity of the Fermi-liquid approach, we only need the value of the vertex function at \( p = p_c \).

This allows us to perform a partial momentum integration in the Dyson equation and write down the equation for the on-shell (\( p = p_c \)) vertex (see Appendices A and C). It reads
\[
J_{ARR} (\mathbf{m}_0) = \sigma + \frac{n_{imp} u_0^2 \varepsilon^2}{4\pi v \delta^2} \int \frac{d\Omega}{4\pi} (1 + \mathbf{n}' \mathbf{n}) J_{ARR} (\mathbf{n}_0) (1 + \mathbf{n}' \mathbf{n}) g_{p,(n-n')} (n,n_0),
\]
(25)
where we denoted \( J_{ARR} (\varepsilon, \mathbf{n}_c) \equiv J_{ARR} (\mathbf{n}_0) \), meaning that the angular dependence of \( J_{ARR} (\varepsilon, \mathbf{n}_c) \) is clearly defined just by the angle between the anisotropy axis and momentum direction (see Fig. 6 illustrating the scattering angles). The energy argument in the scattering times is suppressed. In the isotropic case, equation (25) allows for a simple solution. Since \( J \) is a renormalized current operator, we look for a solution in the form of the polar vector ansatz
\[
J_{ARR} (\varepsilon, \mathbf{n}) = J_1 (\varepsilon) \mathbf{\sigma} + J_2 (\varepsilon) \mathbf{n} + J_3 (\varepsilon) \mathbf{n} (\mathbf{n} \mathbf{\sigma}).
\]
(26)
The ansatz (26) turns Dyson equation (25) into an algebraic one. Changing the integration measure in the expression (13) for the conductivity \( p^2 dp = \varepsilon^2 dp/v^2 \), we arrive at the following suitable formula for the conductivity tensor
\[
\sigma_{\alpha\beta} = \frac{e^2}{6\pi v} \delta_{\alpha\beta} \int \frac{\varepsilon^2 dp}{2\pi \varepsilon} \frac{\varepsilon}{2T} \frac{J_1 (\varepsilon) + J_2 (\varepsilon) + J_3 (\varepsilon)}{\tau_{\varepsilon} + \tau_{\varepsilon}^2}.
\]
(27)
The \( J_1 + J_2 + J_3 \) term is extracted with the help of Eqs. (25) and (20)
\[
J_1 + J_2 + J_3 = \frac{1}{\tau} + \frac{1}{\tau_2}.
\]
(28)

The conductivity then takes the typical Fermi-liquid form
\[
\sigma_{\alpha\beta} = \frac{e^2}{6\pi v} \delta_{\alpha\beta} \int \frac{\varepsilon^2 dp}{2\pi \varepsilon} \frac{\varepsilon}{2T} \frac{1}{\tau_{\varepsilon} + \tau_{\varepsilon}^2}.
\]
(29)

Now, we write down the temperature dependence of the conductivity in the cases of the long- and short-range disorder potentials.

C. Short-range potential

In the case \( p_0 \gg \max \{T/v, \varepsilon_F/v\} \), the structure factor \( g_{p,(n-n')} \) depends only slightly on the scattering angle (between the \( \mathbf{n} \) and \( \mathbf{n}' \) directions) as the scattering is nearly isotropic. This means that \( g(p_c (n - n'))/g(0) = 1 \). The transport scattering time is
\[
\frac{1}{\tau_{\varepsilon}} (\varepsilon) = \frac{2}{3\tau} = \frac{2 n_{imp} u_0^2}{3\pi v \delta^2},
\]
(30)
and the conductivity reads
\[
\sigma_{\alpha\beta} = \sigma \delta_{\alpha\beta}, \quad \sigma = \frac{1}{2\pi} \frac{e^2 v^2 p_0^6}{n_{imp} u_0^2}.
\]
(31)
The conductivity is independent of the chemical potential and temperature in the short-range disorder limit.

D. Long-range potential

This case corresponds to \( p_0 \ll \max \{T/v, \varepsilon_F/v\} \). Now, the scattering is strongly anisotropic. Then, \( p_c^2 (n - n')^2 = 2p_c^2 (1 - \cos \theta) \approx p_c^2 \theta^2 \), and
\[
\frac{1}{\tau_{\varepsilon}} (\varepsilon) = \frac{n_{imp} u_0^2 \gamma_1 v}{8\pi} \frac{1}{p_0^2 \varepsilon^2},
\]
(32)
where \( g_1 = \int_0^\infty g(x) x dx \) (we assume the convergence of the corresponding integral). The conductivity then reads
\[
\sigma_{\alpha\beta} = \frac{8}{3} \frac{e^2}{n_{imp} u_0^2 \gamma_1 v} \frac{\varepsilon_F^2}{v^2} f(\varepsilon_F) \delta_{\alpha\beta},
\]
(33)
\[
f(x) = \left[ 1 + 2\pi^2 x^2 + \frac{7\pi^4}{15} x^4 \right].
\]
In particular, when the doping level is low, \( \varepsilon_F \ll T \) \( x \gg 1) \),
\[
f(x) \approx \frac{7\pi^4}{15} x^4.
\]
(34)
we have
\[
\sigma_{\alpha\beta} \approx \frac{56}{45} \frac{e^2 p_0^2}{n_{imp} u_0^2 \gamma_1 v^2} T^4 \delta_{\alpha\beta}.
\]
(35)
The temperature dependence \( \sigma \sim T^4 \) for long-range disorder of arbitrary form can be qualitatively explained.
in the following manner. The Born transport scattering rate is \( \tau^{-1}_\text{tr} \propto (p_0 \varepsilon)^{-2} \) and the density of states satisfies \( \nu \propto \varepsilon^2 \). For \( T \gg \varepsilon_F \), the characteristic energy of the charge carriers becomes \( \varepsilon \sim T \). Therefore, the Drude conductivity takes the form

\[
\sigma \propto \tau^{-1}_\text{tr}(T) \nu(T) \propto T^4.
\]  

Formula \( (33) \) surprisingly gives the same dependence even for Coulomb disorder. As was pointed out in Refs. \( 26,31 \), the screening length for the Coulomb disorder potential is temperature-dependent and \( p_0 \sim T \). The potential amplitude, however, also depends on temperature \( u_0 \sim c^2 / r \sim c^2 p_0 \propto T \) and the \( T^4 \) dependence survives.

Next, our attention turns to the anisotropic case. As we will see, the longitudinal and transverse conductivities exhibit different temperature dependence. Also, the uniaxial anisotropy introduces additional geometric factors, which are usually needed by experimentalists. In some cases, we managed to obtain exact results.

### IV. ANISOTROPIC CASE

#### A. The Dyson equation and conductivity

For convenience, we now introduce a modified vertex function \( I = I_{\text{ARR}}(2/\tau - 1/\tau) \) and denote \( I_{\text{ARR}}(\varepsilon, n p_0) \equiv I(n n_0) \)

\[
I(n n_0) \left( \frac{2}{\tau(n n_0)} - \frac{1}{\tau(n n_0)} \right) = \sigma \frac{n_{\text{imp}} n_0^2 \varepsilon^2}{4 v^2_\parallel p_0^2 \pi \xi^2} \int \frac{d^4q}{4 \pi} (1 + n' \sigma) I(n' n_0) (1 + n' \sigma) |g_{p_0(n - n'), p_0}|^2.
\]  

The solution to the Dyson equation is sought in the form of the most general polar vector composed of \( \sigma, n \) and \( n_0 \):

\[
I = I_1 \sigma + I_2 n + I_3 n(\sigma n) + I_4 n_0(\sigma n) + I_5 n(n_0 \sigma) + I_6 n_0 + I_7 n_0(n_0 \sigma).
\]  

Plugging \( (38) \) into expression \( (15) \) for the conductivity tensor, we obtain

\[
\sigma_{\alpha\beta}(\varepsilon) = \frac{\varepsilon^2 \xi^2}{2 \pi^2 v_\parallel^2} \int \frac{d\Omega}{4 \pi} \int \left[ \Phi(n n_0) n_\alpha n_\beta + \frac{1}{2} \Psi(n n_0) (n_\alpha n_\beta + n_\beta n_\alpha) \right],
\]  

where \( \Phi = I_1 + I_2 + I_3 + I_5 x, \Psi = I_4 + I_6 + I_7 x, \) and \( x = \cos \theta \).

#### B. Short-range potential

##### 1. \( \delta \)-correlated potential

A weak momentum dependence of the potential leaves it essentially isotropic, because it does not depend on the momentum. Therefore, the answer for the \( \delta \)-correlated potential (its characteristic momentum is \( p_0 = \infty \)) is obtained immediately

\[
\sigma_\parallel = \sigma, \quad \sigma_\perp = \xi^2 \sigma,
\]  

where the conductivity \( \sigma \) is defined in \( (31) \) with \( v = v_\parallel \).

##### 2. Finite-range potential corrections

Now it is clear that the anisotropy in the rescaled basis enters the conductivity expression only as a max\( \{\varepsilon_F, T\} / p_0 \) correction. The first correction in the max\( \{\varepsilon_F, T\} / p_0 \) series is easy to compute, expanding the Dyson equation. The spectrum also acquires anisotropic corrections. They are proportional to the disorder potential amplitude and are irrelevant for sufficiently weak potential. We expand the disorder form factor \( (7) \) according to

\[
g(x) \approx 1 + \delta g(0) x, \quad x = \frac{p^2}{p_0^2} \ll 1.
\]  

Using Eq. \( (39) \) and solving the Dyson equation \( (37) \) with the ansatz \( (38) \), we obtain the corresponding correction to the conductivity (the details of the computation are
hereafter we can consider small scattering angles where

\[ \sigma_{\parallel} = \sigma \left[ 1 - \partial g(0) \frac{1 + 4\epsilon^2 - \pi^2 T^2/3 + \epsilon^2}{5 (v_n p_0)^2} \right], \]

\[ \sigma_{\perp} = \sigma \epsilon^2 \left[ 1 - \partial g(0) \frac{7 + 8\epsilon^2 - \pi^2 T^2/3 + \epsilon^2}{15 (v_n p_0)^2} \right]. \]  (42)

It is worthwhile to note that the anisotropy manifests itself in the temperature-dependent conductivity ratio

\[ \frac{\sigma_{\parallel} \epsilon^2}{\sigma_{\perp}} = 1 - \frac{4}{15} (1 + \epsilon^{-2}) \partial g(0) \frac{\pi^2 T^2/3 + \epsilon^2}{(v_n p_0)^2}. \]  (43)

Equation (43) is one of the central results of the paper. We have just found that the conductivity components in the parallel and transverse directions exhibit different temperature behavior. Although Eq. (43) is obtained in the limit \( p_0 \ll \max\{T/v, \epsilon_F/v\} \), we guess that the temperature dependence of the \( \sigma_{\parallel} \epsilon^2 / \sigma_{\perp} \) ratio should become more pronounced when \( p_0 \sim \max\{T/v, \epsilon_F/v\} \).

**C. Long-range potential**

The long-range potential \( u(p) \) corresponds to the case \( p_0 \ll \max\{\epsilon_F/v_{\parallel}, T/v_{\parallel}\} \). Now the ratio

\[ \kappa_{\epsilon} = \frac{p_0}{p_\epsilon} \ll 1 \]  (44)

becomes an additional small parameter of the problem. The disorder potential does not change much the momentum of an incoming particle \( \delta p \sim p_0 \ll p_\epsilon \). Therefore, hereafter we can consider small scattering angles wherever necessary (see Fig. 7).

The calculations are rather cumbersome. They are presented in Appendix F. The Dyson equation is turned into two coupled differential equations

\[ \Phi''(1 - x^2) + x \Phi'(-3 - \xi^2 + x^2(\xi^2 - 1)) \]

\[ -\Phi(1 + \xi^2 - x^2(\xi^2 - 1)) = \frac{1}{2} (\xi^2 - (\xi^2 - 1)x^2)^{3/2}, \]

\[ \Psi''(1 - x^2) - x[(\xi^2 + 1) - x^2(\xi^2 - 1)] \Psi' = -2\Phi', \]  (45)

where, for brevity, we switched to dimensionless functions \( \frac{\nu^2 \xi^2}{4 v_0^2 g(t, \psi)} \rightarrow (\psi, \psi) \) and \( g_0 = \int_0^\infty g(x) dx \). Naturally, the problem of boundary conditions immediately comes on stage. As will be shown below, it is possible to circumvent it in a number of important limiting cases. The Dyson equation for \( \psi(t) \) determines the \( \psi(t) \) function up to an arbitrary constant. This, however, does not cause any difficulty for the conductivity, because \( \psi(n_{\parallel}) \) enters the integral (39) with \( n \) as a multiplier and forms an odd function of the polar angle.

The resultant expressions for \( \sigma_{\parallel} \) and \( \sigma_{\perp} \) read

\[ \sigma_{\parallel}(T) = \sigma_{1} f \left( \frac{T}{\epsilon_F} \right) \int_{-1}^{1} [\Phi(x) x^2 + \psi(x)] dx, \]

\[ \sigma_{\perp}(T) = \frac{c^2}{2} \sigma_{1} f \left( \frac{T}{\epsilon_F} \right) \int_{-1}^{1} \Phi(x) (1 - x^2) dx, \]  (46)

\[ \sigma_{1} = \frac{8\epsilon^2}{n_{\text{imp}} g t v_0^2 v_{\parallel}^2}, \]

where the function \( f(x) \) is defined in (43).

Next, we explore the geometrical implications of anisotropy in the three limiting cases.

**D. Limiting cases of anisotropy**

1. **Easy plane, \( \xi \gg 1 \)**

The first one is the case of strong anisotropy, when \( \xi \gg 1 \) \( (v_\perp \gg v_{\parallel}) \). Then, the Dyson equations (45) reduce to the first-order differential equation

\[ (\Phi(x))' = \frac{\xi}{2} \sqrt{1 - x^2}, \]

\[ \psi \sim \text{const} + O\left( \frac{1}{\xi} \right). \]  (47)

The constant in \( \psi(x) \) is irrelevant as was mentioned earlier; while the solution, which is analytic in the interval \( x \in [-1, 1] \), reads

\[ \Phi(x) = \frac{\xi}{4} \left( \sqrt{1 - x^2} + \arcsin x \frac{x}{x} \right). \]  (48)

The conductivity can be then written as

\[ \sigma_{\parallel}(T) = \frac{3n^2 \pi}{32} \sigma_{1} f \left( \frac{T}{\epsilon_F} \right), \]

\[ \sigma_{\perp}(T) = \frac{n \pi^2}{32} \sigma_{1} f \left( \frac{T}{\epsilon_F} \right) \left( \frac{1}{2} + 4 \ln 2 \right). \]  (49)
The solution analytic in $\Phi(\xi)$ reads

$$
\Phi(x) = \frac{3}{2} \xi^2 + O(\xi^{-1}), \quad \xi \to \infty. \quad (50)
$$

2. Weak anisotropy, $\xi \approx 1$

Now we turn to the case of $v_\perp \approx v_\parallel$. Let $\xi = 1 + \delta\xi$, $\delta\xi \ll 1$. We expand the vertex function according to $\Phi = \frac{1}{4} + \Phi(x)$ (the value 1/4 corresponds to the isotropic scattering). The corresponding equation reads

$$
\delta\Phi''(1 - x^2) - 4x\delta\Phi' - 2\delta\Phi = -\delta\xi(1 - x^2). \quad (51)
$$

The solution analytic in $x \in [-1, 1]$ has the form

$$
\Phi(x) = \frac{1}{4} \frac{\delta\xi}{12} (x^2 - 5),
\Psi(x) = -(\delta\xi) \frac{x}{6}. \quad (52)
$$

Then, the conductivity reads

$$
\sigma_\parallel(T) = \frac{1}{6} \sigma_1 f \left( \frac{T}{\varepsilon_F} \right) \left( 1 + \frac{4}{15} \delta\xi \right),
\sigma_\perp(T) = \frac{1}{6} \sigma_1 f \left( \frac{T}{\varepsilon_F} \right) \left( 1 + \frac{34}{15} \delta\xi \right). \quad (53)
$$

It is worth noting that to the first order in the deviation from anisotropy, the ratio

$$
\frac{\sigma_\parallel^2}{\sigma_\perp} = 1 + O(\delta\xi^2), \quad \delta\xi \ll 1. \quad (54)
$$

does not exhibit any shift.

3. Easy axis, $\xi \ll 1$

In this case, we are unable to obtain exact coefficients. However, it is enough to say that at $\xi = 0$ there exist finite solutions $\Phi(x)$ and $\Psi(x)$, which lead to some finite integrals with $x^2$ and $x$ in the $[-1, 1]$ range. Then, we have a qualitative answer

$$
\sigma_\parallel = \sigma_1 f \left( \frac{T}{\varepsilon_F} \right) c_1,
\sigma_\perp = \sigma_1 \xi^2 f \left( \frac{T}{\varepsilon_F} \right) c_2, \quad (55)
$$

where $c_1$ and $c_2$ are constants of the order of unity. Then, the conductivity ratio takes the form

$$
\frac{\sigma_\parallel^2}{\sigma_\perp} = \frac{c_1}{c_2} + O(\xi^2), \quad \xi \ll 1. \quad (56)
$$

The above results are summarized in Fig. 8.
port phenomena in 3D systems with Dirac points in their energy spectrum.

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Appendix A: Expression for the polarization operator

In order to work out the polarization-operator diagram in Fig. 3 (left or right), we follow the scheme proposed by Eliashberg. To build an analytical continuation of the expression for the Matsubara polarization operator, we need to establish the analytical properties of the vertex function $J_\alpha(z, z + i\omega_n, \omega_n; p)$ in the whole domain of the complex variable $z$. This is done via the Lehman representation and it was discussed in detail in, e.g., Ref. 13.

The conclusion is that the domain of analyticity of $J_\alpha(z, z + i\omega_n, \omega_n; p)$ is a complex plane with two horizontal cuts: Im$(z + i\omega) = 0$ and Im$(z) = 0$. Since we need a \textit{retarded} vertex function, we put $\omega_n > 0$. Next, the three vertex functions are defined in accordance with the structure of the cuts

$$J_{\alpha}^{RR}(z, z + i\omega, i\omega) \quad \text{if} \quad \text{Im} z > 0,$$
$$J_{\alpha}^{AR}(z, z + i\omega, i\omega) \quad \text{if} \quad -\omega_n < \text{Im} z < 0, \quad (A1)$$
$$J_{\alpha}^{AA}(z, z + i\omega, i\omega) \quad \text{if} \quad \text{Im} z < -\omega_n.$$

The general expression for $\Pi_{\alpha\beta}(i\omega_n)$ then becomes

$$\Pi_{\alpha\beta}(i\omega_n) = -T \sum_{\varepsilon_k} J_{\alpha}(\varepsilon_k, \varepsilon_k + i\omega_n, \omega_n; p) G(\varepsilon_k + i\omega_n, p) j_{\beta} G(\varepsilon_k, p)$$
$$= -\int_C \frac{d\varepsilon}{4\pi i} \tanh \frac{\varepsilon}{2T} J_{\alpha}(\varepsilon, \varepsilon + i\omega_n, \omega_n; p) G(\varepsilon + i\omega_n, p) j_{\beta} G(\varepsilon, p). \quad (A2)$$

The contour $C$ is shown in Fig. 9. As usual, the integral over the large circle vanishes and we are left with integrals over different branches

$$\Pi_{\alpha\beta}(i\omega_n) = \int_{-\infty}^{\infty} \frac{d\varepsilon}{4\pi i} \tanh \frac{\varepsilon}{2T} \bigg\{ J_{\alpha}^{RR}(\varepsilon, \varepsilon + i\omega_n, \omega_n; p) G(\varepsilon + i\omega_n, p) j_{\beta} G(\varepsilon, p) -$$
$$- J_{\alpha}^{AR}(\varepsilon, \varepsilon + i\omega_n, \omega_n; p) G(\varepsilon + i\omega_n, p) j_{\beta} G(\varepsilon, p) + J_{\alpha}^{RR}(\varepsilon - i\omega_n, \varepsilon; p) G(\varepsilon - i\omega_n, p) j_{\beta} G(\varepsilon, p) -$$
$$- J_{\alpha}^{AR}(\varepsilon - i\omega_n, \varepsilon; p) G(\varepsilon, p) j_{\beta} G(\varepsilon - i\omega_n, p) \bigg\}. \quad (A3)$$

Making the analytic continuation $i\omega_n \rightarrow \omega + i0$, we obtain

$$\Pi_{\alpha\beta}^R(\omega) = -v_0^2 \text{tr} \int \frac{d\varepsilon}{4\pi i} \frac{d\varepsilon}{(2\pi)^2} \bigg\{ G(\varepsilon, p) J_{\alpha}^{RR}(\varepsilon, \varepsilon + \omega, \omega; p) G(\varepsilon + \omega, p) j_{\beta} \sqrt{\frac{\tan \frac{\varepsilon + \omega}{2T} - \frac{\varepsilon}{2T}}{2T}} \bigg\} \quad (A4)$$
$$+ G(\varepsilon, p) J_{\alpha}^{RR}(\varepsilon, \varepsilon + \omega, \omega; p) G(\varepsilon + \omega, p) j_{\beta} \sqrt{\frac{\tan \frac{\varepsilon + \omega}{2T} - \frac{\varepsilon}{2T}}{2T}} \bigg\}.$$

Here, the vertex functions $J_{\alpha}$ are defined diagrammatically in Fig. 3c. They obey the Dyson equation presented in a diagrammatic form in Fig. 3d. The disorder is static and does not cause a change of frequency in the diagrammatic loops. Thus, the Dyson equation takes an especially elegant form

$$J_{\alpha}(i\varepsilon_k, \varepsilon_k + i\omega_n, \omega_n; q) = \sigma_\alpha + \frac{n_i m v_F^2}{p_0} \int \frac{dp}{(2\pi)^3} J_{\alpha}(i\varepsilon_k, p) J_{\alpha}(i\varepsilon_k + i\omega_n, \omega_n; q) G(i\varepsilon_k + i\omega_n, p). \quad (A5)$$
We are interested in the zero-frequency response of the system and set the external frequency $\omega = 0$. After the analytic continuation, the vertex function is split into $RRR, AAR$ and $ARR$ parts

$$J_\alpha^{ARR}(\varepsilon, q) = \sigma_\alpha + \frac{n_{imp}b_0^2}{p_0^2\varepsilon^2} \int \frac{dp}{(2\pi)^3} g_{q-p,n_0} G^A(\varepsilon, p) J_\alpha^{ARR}(\varepsilon, p) G^R(\varepsilon, p),$$

$$J_\alpha^{RRR}(\varepsilon, q) = \sigma_\alpha + \frac{n_{imp}b_0^2}{p_0^2\varepsilon^2} \int \frac{dp}{(2\pi)^3} g_{q-p,n_0} G^R(\varepsilon, p) J_\alpha^{RRR}(\varepsilon, p) G^R(\varepsilon, p),$$  

$$J_\alpha^{AAR}(\varepsilon, q) = \sigma_\alpha + \frac{n_{imp}b_0^2}{p_0^2\varepsilon^2} + \int \frac{dp}{(2\pi)^3} g_{q-p,n_0} G^A(\varepsilon, p) J_\alpha^{AAR}(\varepsilon, p) G^A(\varepsilon, p).$$

The learned reader already knows that only $J^{ARR}$ (so called singular) vertex undergoes a strong (non-perturbative) renormalization due to the interaction with impurities, while the other two exhibit weak perturbative corrections. The reason is, of course, the position of the poles of the Green’s functions. Since we claim some rigor, we discuss this issue in detail in Appendix B.

**Appendix B: One-loop structure of the Green’s function**

1. **General expression**

The general expression for the one-loop self-energy reads

$$\Sigma^R(\varepsilon, q) = \frac{n_{imp}b_0^2}{p_0^2\varepsilon^2} \int \frac{dp}{(2\pi)^3} (\varepsilon + v_n p_\sigma - v_n^2 p_\sigma^2) g_{q-p,n_0} \frac{d\Omega}{4\pi},$$

and its imaginary part

$$\text{Im} \Sigma^R(\varepsilon, q) = -\frac{n_{imp}b_0^2\varepsilon^2}{4\pi v_n^2 p_0^2\varepsilon^2} \int (1 + n_0 \sigma) g_{q-p,n_0} \frac{d\Omega}{4\pi}.$$  

(B1)

(B2)

The self energy has the following tensor form

$$\Sigma^R(\varepsilon, q) = \alpha(\varepsilon, q) + \beta(\varepsilon, q)\sigma - \frac{i}{2\tau(\varepsilon, q)} \frac{i\sigma s(\varepsilon, q)}{2\tau(\varepsilon, q)},$$

(B3)

where $s$ and $\beta$ are vectors in the $(q, n_0)$ plane and

$$\alpha = \frac{|\beta|}{\varepsilon} \frac{2}{\varepsilon} \min\{p_0 v_n, \varepsilon\} \frac{1}{\varepsilon^2} \ll 1.$$  

(B4)

For the case of isotropic potential, the anisotropic short range potential $(p_0 \gg p_s)$ and anisotropic long range potential $(p_0 \ll p_s)$, vector $s \equiv n$ and the scattering rates $\tau$ and $\tau_1$ are immediately extracted and presented in (B3).

As is seen from the structure of the integrals (B1) and (B2),

$$\tau(\varepsilon, -q, \theta) = \tau(\varepsilon, q, \theta);$$

$$s(\varepsilon, -q, \theta) = -s(\varepsilon, q, \theta) \frac{\tau_1(\varepsilon, -q, \theta)}{\tau_1(\varepsilon, q, \theta)}.$$  

(B5)

Here $\theta$ is the angle between $q$ and $n_0$ and the parity relation follows from the measure invariance under the change $\theta \rightarrow \pi - \theta, \varphi \rightarrow \varphi + \pi$.

Now we may write the Green’s function in the standard way

$$G^R(\varepsilon, q) = \frac{\varepsilon - \alpha + \frac{i}{2\tau_1}}{\varepsilon - \alpha + \frac{i}{2\tau_1}} + \frac{\sigma(q + \beta - \frac{is}{\varepsilon})}{(\varepsilon - \alpha + \frac{i}{2\tau_1}) - (q + \beta - \frac{is}{\varepsilon})^2}.$$  

(B6)

In a typical computation, we need to integrate products of the type $G^R(\varepsilon, q) G^R(\varepsilon, q)$, $G^R(\varepsilon, q) G^A(\varepsilon, q)$ and $G^A(\varepsilon, q) G^R(\varepsilon, q)$ over the momentum $q \in (-\infty, \infty)$. Therefore, we need to know the position of the poles of the Green’s function in the $q$-domain. The retarded function $G^R(\varepsilon, q)$ obeys the causality condition, which means that its poles, defined by the equation

$$\det[\varepsilon - v_n q_\sigma - \Sigma^R(\varepsilon, q)] = 0,$$  

(B7)

lie in the lower half-plane of the complex variable $\varepsilon$, i.e.

$$\frac{1}{\tau(\varepsilon, q, \theta)} + \frac{\sigma s(\varepsilon, q, \theta)}{\tau_1(\varepsilon, q, \theta)} > 0,$$  

(B8)

for any $\varepsilon$ and $\theta$; $n$ stands for $q/q$. Since we need the position of the poles in the $q$-plane, we solve the spectral
Now, we see that while integrating the product \( G^A(\varepsilon,q) \) in the complex plane \( q \) is always such as depicted in Fig. 11(a). The poles of \( G^G(\varepsilon,q)G^H(\varepsilon,q) \) and \( G^A(\varepsilon,q)G^R(\varepsilon,q) \) are sketched in Figs. 11(b) and 11(c) respectively. Now, we see that while integrating the product \( G^G G^R \) or \( G^A G^A \) over \( q \) the contour of integration can be deformed to pass far from the poles. When integrating the product \( G^A G^R \), the contour is squeezed between the nearby poles and the deformation is no longer possible, leading to their enhanced contribution.

**Appendix C: Dyson equation**

1. **Structure of \( J^{ARR}(\varepsilon,p) \)**

In order to find the conductivity, we need to solve the Dyson equation for the vertex function \( J^{ARR}(\varepsilon,p) \)

\[
J_1(q)\sigma + J_2(q)\frac{p_q q}{q^2} + J_3(q)\frac{q(q\sigma)}{q^2} = \sigma + \frac{\hbar v_{\text{imp}}^2}{p_0^2} \nonumber
\]

\[
\times \int \frac{dp}{(2\pi)^3 q_l q_l - p} J_1(p) [p_0^2 - p^2] + J_2(p) \frac{p_0 p}{p_0^2} [p_0^2 - p^2 + 2p_0^2 (p\sigma)] + J_3(p) \frac{p_0^2}{p_0^2} [p_0^2 - p^2 (p\sigma) + 2p_0 p^2] \nonumber
\]

For the integral (C2) to converge, the vertices \( J_{1,2,3}(p) \) must be bounded when \( p \to \infty \). As in expression for the conductivity, the integrals has a sharp peak at \( p = p_c \) of width \( \Delta p \sim 1/(v\tau) \) near the Fermi surface. Generally, we cannot restrict the computation of the integral by expanding the integral near the Fermi surface. Such an expansion leads to a significantly deformed integrand (see Fig. 11(a)). First, however we discuss the vertex at \( p = p_c \).

2. **On-shell vertex: \( J^{ARR}(\varepsilon,p_c) \)**

If we are interested in the value of \( J_{1,2,3}(p) \) at \( p = p_c \), then the range of momentum integration is the sphere of convergence of the disorder structure factor \( T \), namely the sphere of radius \( p_0 \) with the center at the external momentum \( q = p_c n \). There are two distinctive contributions. The first one comes from the integration inside the small sphere: \( |p - p_c n| \leq 1/(\tau v) \), where the integrand is peaked due to the proximity of the poles of \( G^A G^R \); the second one comes from a thick shell \( \tau_{1/2}(\varepsilon) \lesssim |p - p_c n| \lesssim p_0 \). Alternatively, one can think of the first contribution as the one coming from the singularities of \( G^A G^A \) and the second of the singularities of \( g_{n p_c - p} \). Simple estimates give

\[
\int \frac{dp}{p_0} G^A(p) J_0 G^R(p) g_{n p_c - p} \sim \tau_{1/2} p_0 \nonumber
\]

\[
\int \frac{dp}{p_0} G^A(p) J_0 G^R(p) g_{n p_c - p} \sim \tau_{1/2} p_0 \nonumber
\]

Comparing the two contributions, we see that if the disorder cut-off \( p_0 \) is not too high, the contribution from the thick shell is \( 1/(\tau v) \times (p_0/p_c)^3 \) smaller than the one from the small sphere due to the \( G^A G^A \) product. Therefore, one can indeed discard the contribution from the shell \( |p - p_c n| \lesssim p_0 \) (or equivalently, the contrib-
the leading order of the \( \varepsilon \cdot \varepsilon \)-expansion, one can take into account only the poles of the \( G^{R}G^{A} \) term.

3. Vertex \( J^{ARR}(\varepsilon,p) \) at \( p \rightarrow \infty \)

The same logic of estimates holds for the large \( p \) behavior of the vertex. We now plug ansatz (27) into the Dyson equation (A6) and employ the estimates (C3), modified for the external momentum \( q \gg p_{c}, p_{0} \). The computation is simple though tedious. Solving the system of self-consistent equations, one arrives at the following asymptotics

\[
\begin{align*}
J_{1}(q) & = 1 + \beta_{1} g_{q} p_{c}^{2} J_{0}, \\
J_{2}(q) & = \frac{\beta_{2} q}{p_{c}} p_{0} \partial g_{q} p_{c}^{2} J_{0}, \\
J_{3}(q) & = \frac{\beta_{3} g_{q} p_{c}^{2}}{p_{0}} J,
\end{align*}
\]

where \( J \equiv J_{1}(p_{c}) + J_{2}(p_{c}) + J_{3}(p_{c}) \) and \( \beta_{1,2,3} \) are constants of the order of unity. We see that the estimate for the vertex in the isotropic case can be rewritten as

\[
J(q) - \sigma \sim v_{g}(q),
\]

where \( r \) is some restricted vector function of the momentum \( q \). Therefore, the renormalized current vertex \( J(q) \) approaches the bare one \( \sigma \) at large momenta \( q \) with the rate proportional to the structure factor \( g(q) \).

4. On shell Dyson equation

Now we are able to write down the Dyson equation in a somewhat simplified form performing integration over the absolute value of the momentum. As was argued previously, the most important contribution comes from the peak in the \( G^{R}G^{A} \) product at \( p' = p_{c} \). Therefore, we put \( p' \) in the argument of \( J(\varepsilon,pn') \) and \( g'(p - pn', p_{0}) \) inside the integral equation (A6) equal to \( p_{c} \). Moreover, we take into account the fact that in all the limiting cases considered (isotropy, anisotropy with short and long-range disorder) the Green’s function assumes a simplified form (23) rather than (16) (see Appendix F).

Changing \( p^{2} = \zeta, \, dp = v_{g}(d\zeta)/2\zeta \), and changing the lower limit of integration to \(-\infty \), we perform the integration

\[
\int_{0}^{\infty} \frac{p^{2} dp}{\left( \left( \varepsilon^{2} - v_{g}^{2}(p - \frac{1}{2\xi})^{2} \right) - v_{g}^{2}(p + \frac{1}{2\xi})^{2} \right)^{2}} \approx \frac{\pi}{2v_{g}^{2} \frac{1}{\xi} + \frac{1}{\tau_{1}}},
\]

This way we arrive at the Dyson equation (23) and (24) for the modified vertex.

Appendix D: Estimate of the regular term in conductivity

Thanks to the factor \( \varepsilon^{2} - p^{2} \) in the nominator of (16) we can substitute

\[
(p^{2} - \varepsilon^{2}) G^{A}(\varepsilon, p) \approx (\varepsilon + p \sigma) \left[ 1 + O\left( \frac{1}{\varepsilon \tau} \right) \right],
\]

and the contour of integration is no longer squeezed between the proximate poles of the product \( G^{R}G^{A} \). The path therefore can be deformed in any suitable way. Estimating at \( p \sim \max\{p_{c}, p_{0}\} \), \( G^{R}G^{A} \sim 1/\varepsilon^{2} \), we obtain

\[
I \sim \delta_{\alpha \beta} \left| \max\{p_{c}, |J(\varepsilon, p_{c}) - \sigma|, p_{0}|J(\varepsilon, p_{0}) - \sigma| \right) \frac{1}{\varepsilon^{2}}.
\]

On the other hand, the convergent expression \( \Pi \) is completely determined by the behavior of \( J_{\alpha}^{ARR}(\varepsilon, p) \) in the vicinity of the peak and one can substitute \( J_{\alpha}(\varepsilon, p) \) with \( J_{\alpha}(\varepsilon p_{F}, p_{F}) \). The integral is then estimated as

\[
\Pi \sim \delta_{\alpha \beta} J(\varepsilon, p_{c}) - \sigma \frac{\varepsilon^{2} \tau}{p_{c}^{3}}.
\]

Combining Eqs. (D3) and (C5), we obtain relation (17).
Appendix E: Conductivity, short-range potential

In the $\delta$-correlated (momentum-independent) case, the vertex functions $\Psi$ and $\Phi$ are angle-independent and, as follows from Eq. (25)

$$
\Phi(n_0 n) \equiv \Phi_0 = \frac{3\tau}{2}, \quad \Psi(n_0 n) = 0. \tag{E1}
$$

In the case of small momentum-dependent correction to form factor [11], these also acquire small angle-dependent corrections

$$
\Phi(n_0 n) = \Phi_0 + \delta \Phi(n_0 n), \quad \Psi(n_0 n) = \delta \Phi(n_0 n) \tag{E2}
$$

of the order of $(p/\mu)^2$. Thus, the formula for the conductivity (see Eq. (39)) becomes

$$
\sigma_{\alpha\beta} = \sigma \times \left( \delta_{\alpha\beta} + \frac{\delta \Phi_0 - \delta \Phi_2}{4\tau} \delta_{\alpha\beta} + \frac{3 \delta \Phi_2 - \delta \Phi_0 + 2 \delta \Psi_1}{4\tau} n_{\alpha\beta} n_{\alpha\beta} \right), \tag{E3}
$$

where $(\delta \Psi_1, \delta \Phi_0) = \int_1^{\frac{1}{2}} (\delta \Psi, \delta \Phi) x^2 dx$. Now $\delta \Phi_0$ and $\delta \Psi_1$ are easily extracted perturbatively from the Dyson equation (see the next Appendix for details). The corrections can be written as

$$
\delta \sigma_\perp = \xi^2 \sigma \frac{\delta \Phi_0 - \delta \Phi_2}{4\tau},
$$

$$
\delta \sigma_\parallel = \sigma \frac{\delta \Phi_2 + \delta \Psi_1}{2\tau}. \tag{E4}
$$

We thus obtain the conductivity as a function of the energy

$$
\delta \sigma_\perp = -\sigma \frac{\xi^2}{5} \frac{p_\perp^2}{p_0^2} \frac{\partial g}{g} (1 + 4\xi^{-2}),
$$

$$
\delta \sigma_\parallel = -\sigma \frac{\xi^2}{15} \frac{p_\perp^2}{p_0^2} \frac{\partial g}{g} (7 + 8\xi^{-2}). \tag{E5}
$$

Plugging these into (44), we determine the temperature dependence of the conductivity [42].

Appendix F: Solution of the anisotropic Dyson equation

1. Short-range potential

We now need to compute the anisotropic corrections to the spherically-symmetric self-energy. From the structure of (12), we see that we need to compute the following averages

$$
n_{\text{imp}} = \frac{\tau}{\mu^2 \pi} \int \frac{d\Omega'}{4\pi} g'_{\nu_\nu}(n-n') n_0 n \tag{F1}
$$

$$
= -\frac{\partial g(0)p^2}{3\tau p_0^2} [(1 - \xi^{-2})(n_0 n) + \xi^{-2} n],
$$

$$
nen_{\text{imp}} = \frac{\tau}{\mu^2 \pi} \int \frac{d\Omega'}{4\pi} g'_{\nu_\nu}(n-n') n_0 n_{\alpha\beta} = \frac{1}{6\tau}
$$

$$
\times \left[ (\delta_{\alpha\beta} + 1 + \frac{\partial g(0)p^2}{p_0^2} \left( (1 - \xi^{-2})(n_0 n) + \frac{12 + 9\xi^{-2}}{5} \right) \right]
$$

$$
+ \frac{2\partial g(0)p^2}{5p_0^2} (1 - \xi^{-2})(n_0 n) n_{\alpha\beta} n_{\alpha\beta} \tag{F1}
$$

Therefore, we see that the self-energy retains its isotropic form with the corrections to the scattering times:

$$
\delta \frac{1}{\tau} = \frac{1}{\tau} \frac{\partial g(0)p^2}{p_0^2} \left( (1 - \xi^{-2}) \left( n_0 n \right)^2 + \frac{1}{3} + 2\xi^{-2} \right), \tag{F2}
$$

$$
\delta \frac{1}{\tau_{\text{tr}}} = \frac{1}{\tau_{\text{tr}}} \frac{\partial g(0)p^2}{p_0^2} \left( \frac{5}{3} (1 - \xi^{-2})(n_0 n)^2 + \frac{1 + 7\xi^{-2}}{3} \right). \tag{F3}
$$

Following the same steps as in the isotropic case, we arrive at the following equations for the vertex functions

$$
\delta \Phi(x) = -\frac{2\tau}{15} \frac{\partial g(0)p^2}{p_0^2} (1 + 4\xi^{-2}) + \frac{\delta \Phi_0 - \delta \Phi_2}{4},
$$

$$
\delta \Psi_1(x) = -\frac{8\tau}{15} \frac{\partial g(0)p^2}{p_0^2} (1 - \xi^{-2}) x + \frac{\delta \Psi_1 + \delta \Psi_1}{2}. \tag{F4}
$$

Solving Eqs. (F3) in a self-consistent way, we find

$$
\delta \Phi_0 = -\frac{6}{5} \tau (1 + 4\xi^{-2}) \frac{\partial g(0)p^2}{p_0^2},
$$

$$
\delta \Phi_2 = \frac{2}{5} \tau (1 + 4\xi^{-2}) \frac{\partial g(0)p^2}{p_0^2}, \tag{F4}
$$

$$
\delta \Psi_1 = -\frac{8}{15} \tau (1 - \xi^{-2}) \frac{\partial g(0)p^2}{p_0^2}. \tag{F4}
$$

From the last equations, we easily recover Eqs. (5).

2. Long-range disorder

Expanding in the scattering angle, we have

$$(n - n') n_0 = (1 - \cos \theta') \cos \theta + \sin \theta' \sin \theta \cos \varphi \approx \theta' \sin \theta \cos \varphi, \tag{F5}
$$

$$(n - n')^2 = 2 - 2 \cos \theta \approx \theta'^2, \quad \cos \theta' \sin \varphi \approx \theta' \sin \varphi \sin \varphi.$$
Next, we extend the upper limit of integration over $\theta'$ to $+\infty$, and we obtain

$$\text{Im} \Sigma(\varepsilon, p_n) = -\frac{n_{\text{imp}} e^2 u_0^2}{4 \pi v^2 |p_0|^{2\varepsilon}} \int_0^{2\pi} \frac{d\varphi}{2\pi} \left( 1 + \frac{1}{1 - \frac{v^2}{c^2}} \right) \left( 1 + \frac{1 + \sigma_\varepsilon \theta' \cos \varphi}{1 - \sigma_\varepsilon \theta' \sin \varphi} \right) \int_0^\infty \frac{\theta'}{2} (1 - \xi^2)^2 \sin^2 \theta \cos^2 \varphi + \xi^2 \right)$$

$$= -\frac{1}{2\tau(m_0)} \frac{\sigma_n}{\tau(\tau(\tau(m_0)))}.$$ (F6)

In addition to (24), we will need one more scattering time

$$\frac{1}{\tau_{\text{tr}}(\varepsilon)} = \frac{n_{\text{imp}} e^2 u_0^2}{2\pi v^2 |p_0|^{2\varepsilon}} \int d\Omega \frac{d\theta}{4\pi} g_p \eta_{n'(n-m),m_0} (1 - \cos \theta_{nm}) \cos^2 \varphi$$

(see Fig. 3 for the definition of the angle $\varphi$). To obtain the explicit formulas for all scattering times, we expand the argument of $g_p \eta_{n'(n-m),m_0}$ and integration domain $d\Omega$.

Introducing the notation

$$\Delta(\theta) \equiv \Delta(m_0) = [(\sigma_\varepsilon^2 - 1) \sin^2 \theta + 1]^{1/2},$$

where $g_0$ is defined after Eq. (45), we obtain results for the cross-sections

$$\frac{1}{\tau(m_0)} = \frac{1}{\tau_0} \Delta(m_0),$$

$$\frac{1}{\tau_{\text{tr}}(\varepsilon)} = \frac{n_{\text{imp}} e^2 u_0^2}{4 \pi v^2 |p_0|^{2\varepsilon}} \int d\Omega g_p \eta_{n'(n-m),m_0} \frac{1}{\Delta^2(m_0)},$$

$$\frac{1}{\tau_{\text{tr}}(\tau(m_0))} = \frac{1}{\tau_{\text{tr}}(\tau(m_0))} \left( \Delta^2(m_0) + 1 \right).$$ (F9)

(Here $g_1$ is defined after Eq. (62)). As one can see from the expression for $\text{Im} \Sigma$ (F6), the Green’s function retains its simple structure (23). One needs to pay particular attention to the fact that the scattering time is now a function of the direction of the incoming particle with respect to the anisotropy axis. Now we plug the Green’s functions (23) and ansatz (38) into the Dyson equation (37).

The most obvious step in solving the Dyson equation (37) is to take the zero order in the $\kappa$-expansion and to set $n' = n$ in the peaked integrand. One immediately arrives at an algebraic equation for all the constants. Despite the fact that one is able to immediately obtain some of the vertices: $I_1 = 1$, $I_5 = I_7 = 0$, we also obtain a meaningless equation $I_2 = I_3 = 1 + I_2$ and the vertices $I_4$, $I_6$ remain undefined. This is a signature of the fact that one needs to take the next term in the $\kappa$-expansion.

The $\kappa$ expansion in turn leads to the emergence of the derivatives of functions $I$ in the structure of the equation.

We use the commutation relation and find $(1 + n\sigma)(1 + n\sigma) = 2(1 + n\sigma)$.

$$(1 + n\sigma)I(1 + n\sigma)$$

$$= 2(I_1 + I_2 + I_3 + I_4 \eta_{n,n_0})n(1 + n\sigma)$$

$$+ 2(I_4 + I_5 + I_7 \eta_{n,n_0})n(1 + n\sigma).$$ (F10)

Whenever necessary, we expand

$$n' = n + \delta n,$$

$$I(n',n_0) = I(n_0 + \delta n_0)$$

$$= I(n_0) + n_0 \delta n \delta n_0 I(n_0) + \frac{(n_0 \delta n)^2}{2} \frac{\partial^2}{\partial n_0^2} I(n_0) + ...$$ (F11)

Next, we perform the integration over angles

$$\int d\Omega ' \frac{d\theta'}{4\pi} g_p \eta_{n'(n-m),m_0} \delta n$$

$$= \frac{n_{\text{imp}} e^2 u_0^2}{4 \pi v^2 |p_0|^{2\varepsilon}} \int d\Omega d\theta$$

$$\frac{n_{\text{imp}} e^2 u_0^2}{4 \pi v^2 |p_0|^{2\varepsilon}} \int d\Omega ' d\theta'$$

$$\frac{n_{\text{imp}} e^2 u_0^2}{4 \pi v^2 |p_0|^{2\varepsilon}} \int d\Omega d\theta$$

$$= \frac{1}{\tau_{\text{tr}}(\tau(m_0))} \left[ \Delta^2 \delta _{\alpha \beta} - (\xi^2 - 1) n_{\alpha,n_0} n_{\beta} \right.$$}

$$- \xi^2 n_{\alpha,n_0} n_{\beta} + (\xi^2 - 1) (n_0 n_{\alpha,n_0} n_{\beta} + n_{\alpha,n_0} n_{\beta}) \right].$$ (F12)

The Dyson equation is split into seven differential equations. In terms of $\Phi(x)$ and $\Psi(x)$, we rewrite it as follows
Summing up the equations, we obtain (F13).

\[
I_1 \left( \frac{2}{\tau} - \frac{1}{\tau_{tr}} \right) = 1 + \frac{2\Delta^2}{\tau_{tr}} \Phi, \\
I_2 \left( \frac{2}{\tau} - \frac{1}{\tau_{tr}} \right) = \Phi \left( \frac{2}{\tau} - \frac{1}{\tau_{tr}} \right) - x\Phi' \left( \frac{2}{\tau_{tr}} \right) + \frac{1}{\tau_{tr}} \Phi''(1 - x^2) + I_5 x, \\
I_3 \left( \frac{2}{\tau} - \frac{1}{\tau_{tr}} \right) = \Phi \left( \frac{2}{\tau} - \frac{2}{\tau_{tr}} - 2\xi^2 \right) - x\Phi' \left( \frac{4}{\tau_{tr}} \right) + \frac{1}{\tau_{tr}} \Phi''(1 - x^2), \\
I_4 \left( \frac{2}{\tau} - \frac{1}{\tau_{tr}} \right) = \frac{2}{\tau_{tr}} (\Phi(\xi^2 - 1)x + \Phi' + \frac{1}{\tau} - \frac{1}{\tau_{tr}}) - x\Phi' \left( \frac{2}{\tau_{tr}} + \frac{1}{\tau_{tr}} \right) + \frac{1}{\tau_{tr}} \Phi''(1 - x^2), \\
I_5 \left( \frac{2}{\tau} - \frac{1}{\tau_{tr}} \right) = \frac{2}{\tau_{tr}} (\Phi(\xi^2 - 1)x + \Phi'), \\
I_6 \left( \frac{2}{\tau} - \frac{1}{\tau_{tr}} \right) = \frac{2}{\tau_{tr}} \Phi + \frac{1}{\tau} - \frac{2}{\tau_{tr}} \Phi' + \frac{1}{\tau_{tr}} \Phi''(1 - x^2), \\
\left( \frac{2}{\tau} - \frac{1}{\tau_{tr}} \right) I_7 = -\frac{2}{\tau_{tr}} \Phi(\xi^2 - 1) + \frac{2}{\tau_{tr}} \Phi'.
\]

Summing up the equations, we obtain (F13).

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