Electronic hydrodynamics and the breakdown of the Wiedemann-Franz and Mott laws in interacting metals

Andrew Lucas\textsuperscript{a} and Sankar Das Sarma\textsuperscript{b}
\textsuperscript{a} Department of Physics, Stanford University, Stanford, CA 94305, USA
\textsuperscript{b} Condensed Matter Theory Center and Joint Quantum Institute, Department of Physics, University of Maryland, College Park, MD 20742 USA

ajlucas@stanford.edu

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Abstract: We present the theory of thermoelectric transport in metals with long-lived quasiparticles, carefully addressing the interplay of electron-electron scattering as well as electron-impurity scattering, but neglecting electron-phonon scattering. In Fermi liquids with a large Fermi surface and weak electron-impurity scattering, we provide universal and simple formulas for the behavior of the thermoelectric conductivities across the ballistic-to-hydrodynamic crossover. In this regime, the electrical conductivity is relatively unchanged by hydrodynamic effects. In contrast, the thermal conductivity can be parametrically smaller than predicted by the Wiedemann-Franz law. A less severe violation of the Mott law arises. We quantitatively compare the violations of the Wiedemann-Franz law arising from (i) momentum-conserving electron-electron scattering in the collision integral, (ii) hydrodynamic modifications of the electron-impurity scattering rate, and (iii) thermal broadening of the Fermi surface, and show that (i) is generally the largest effect. We present simple formulas for electrical and thermal magnetococonductivity across the ballistic-to-hydrodynamic limit, along with a more complicated formula for the thermoelectric magnetococonductivity. In a finite magnetic field, the Lorenz number may be smaller or larger than predicted by the Wiedemann-Franz law, and the crossover between these behaviors is a clear prediction for experiments. The arbitrarily strong violation of the Wiedemann-Franz law found in our work arises entirely from electron-electron interaction effects within the Fermi liquid paradigm, and does not imply any non-Fermi liquid behavior. We predict clear experimental signatures of bulk hydrodynamics in high-mobility 2D GaAs semiconductor structures, where a spectacular failure of the Wiedemann-Franz law should persist down to very low temperatures in high-quality and low-density samples.

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Introduction

One of the most important open problems in condensed matter physics is a non-perturbative understanding of interaction effects on the electronic transport properties of metals. The subject is vast and an enormous literature exists, spanning almost a century. Due to unresolved experimental puzzles such as the high temperature linear-in-temperature resistivity of the normal phase of high-temperature superconductors at optimal doping [1], the study of interaction-limited transport has taken on much significance over the last two decades. Typically, one assumes that strong electron-electron interactions lead to a breakdown of Fermi liquid physics and of a quasiparticle description, such that the resulting metal is a non-Fermi liquid (NFL) [2, 3, 4, 5]. Despite much effort, the understanding of NFLs is very challenging and remains incomplete.

A common heuristic used in experiments to probe the breakdown of Fermi liquid phenomenology is the Wiedemann-Franz (WF) law [6, 7], which states that the ratio of thermal and electrical conductivity is

\[ \frac{\kappa}{\sigma T} \approx \frac{\pi^2 k_B^2}{3 e^2}. \]  

(1)

The ratio on the left hand side above is called the Lorenz number, and the constant on the right hand side is the ideal Lorenz number experimentally observed in most normal elemental metals at room temperatures.
The WF law arises in a conventional metal at temperatures $T \ll T_F$, the Fermi temperature, whenever electron-electron collisions are negligible, and elastic scattering processes dominate [8]. Transport properties of normal metals are dominated by electron-phonon scattering, which is essentially quasi-elastic at room temperatures. The theoretical prediction (1) therefore holds very well for most normal metals [9]. Furthermore, NFLs will usually (but not always) manifest a Lorenz number much smaller than predicted by (1) [3]. In some instances, the implication that bad metals (apparent NFLs with large resistivity [2, 3, 4]) violate the WF law has been “inverted” to conclude that a failure of the WF law by itself is evidence for the existence of a NFL [10]. Of course, this conclusion is not logically necessary: it is possible for Fermi liquids to violate (1) even in the very low temperature limit [11]. In particular, (1) is violated whenever the scattering processes which dominate transport are inelastic.

The purpose of this paper is to – within the Fermi liquid paradigm – describe the interplay of electron-electron interactions and impurities. A careful understanding of this interplay is essential to develop a NFL transport theory. We will see that some of the thermoelectric transport phenomena that occur in NFLs also occur in FLs. Our description of ‘unconventional’ FL transport phenomena both provides a “less strange” setting in which to understand interaction-limited transport physics, and sheds light on which transport phenomena rely on the breakdown of FL theory, and which do not. Both the hydrodynamic FLs which we consider, and the NFLs outlined previously, are strongly interacting electron systems. One main difference is that NFL behavior is commonly believed to emerge due to a putative nearby quantum critical point. The WF violation and related ‘exotic’ transport properties being studied in our current work arise not from any hidden quantum criticality, but from strong interaction effects within the metallic Fermi liquid itself. The hidden quantum criticality which may be responsible for NFL physics will most strongly manifest itself in specific temperature-dependent corrections to the effective electron-impurity scattering rate.\(^1\) It is an interesting question, beyond the scope of the current work, whether an interplay of quantum criticality and hydrodynamic effects coexist in strongly correlated materials; see recent discussion in [13, 14].

Expanding upon earlier works such as [3, 11], we provide a comprehensive and quantitative theory of the finite temperature breakdown of the WF law in metals described by Fermi liquid theory. In particular, we confirm the results of [11] that the Lorenz ratio becomes parametrically smaller than (1) in the hydrodynamic limit [15], where non-umklapp electron-electron scattering processes occur much faster than electron-impurity or electron-phonon processes. This effect is universal and does not rely on any details of the interactions or band structure. Any metal, if driven to a hydrodynamic regime,\(^2\) will necessarily violate (1). This violation does not necessarily imply NFL behavior.

We will also describe the interaction-driven breakdown of the related Mott law for thermoelectric conductivity:

$$\alpha = -\frac{\pi^2 k_B^2 T}{3 e} \frac{\partial \sigma}{\partial \mu}, \hspace{1cm} (2)$$

with $\mu = k_B T_F$ the chemical potential, or Fermi energy. We will see that the Mott law holds in the same parameter regimes as the WF law; however, its breakdown is not as dramatic.

The reason that hydrodynamic breakdowns of the WF and Mott laws are not observed in typical metals, even at very low temperatures, is simply a reflection of the fact that most metals are rather dirty, and exhibit electron-electron scattering rates which are negligibly small compared to electron-impurity scattering rates. Elemental 3D metals are never in the hydrodynamic regime, even when they are relatively pure – at low (high) temperatures, electron-impurity (-phonon) interactions are stronger

\[^1\] At finite temperatures there is generally simply a crossover (not a phase transition) between the quantum critical fan and conventional portions of the phase diagram [12]. The impact of quantum critical fluctuations is most clearly seen in the unconventional temperature dependence of scattering rates.

\[^2\] In the absence of umklapp, this can be done by making the crystal exceptionally pure so as to drastically reduce the electron-impurity scattering rate.
than the corresponding electron-electron interaction. However, effectively 2D metals (e.g. graphene, high-
mobility 2D semiconductor layers) can often be clean enough for the electron-electron scattering rate to
surpass the electron-impurity scattering rate at low to intermediate temperatures (where electron-phonon
scattering is relatively weak). These systems can be in the hydrodynamic regime. We urge experiments in
high-mobility low-density modulation-doped 2D GaAs electron and hole systems [16] in order to verify our
predictions of hydrodynamics-induced breakdowns of the WF and Mott relations arising from electron-
electron interactions. These high-quality GaAs systems, which are routinely used for studies of the
fractional quantum Hall effect, are exceptionally pure and easily enter the hydrodynamic regime in a
0.1–10 K temperature range.

In fact, recent experiments on graphene [17, 18], PdCoO$_2$ [19], WP$_2$ [20] and GaAs [21] have all
observed evidence for the collective hydrodynamic flow of electrons. Interaction-limited transport phe-
nomena in low density oxides such as SrTiO$_3$ [22, 23] may also have a hydrodynamic character. Our results
complement the existing theoretical literature on hydrodynamic transport [24, 25, 26, 27, 28, 29, 30] and
contain new hydrodynamic predictions for experiments in the materials listed above. In graphene, break-
downs of the WF [31] and Mott [32] laws have been reported and attributed to hydrodynamic effects. A
demonstration of quantitative consistency between bulk transport phenomena and more direct probes of
viscous effects such as nonlocal resistance will be strong evidence for the hydrodynamic nature of electron
flow in these systems.

Consistent with our goal of understanding the role of electron-electron interaction in metallic transport
(and particularly, the hydrodynamic regime), we ignore electron-phonon interaction in our work. This
is not because electron-phonon interaction is generically unimportant – as we noted previously, room
temperature transport in normal metals is dominated by phonons – but because the role of electron-
phonon interaction in metallic transport is well-understood [8]. Neglecting phonons keeps our theory
transparent and tractable. It is straightforward to add electron-phonon scattering effects to our theory.

1.1 Outline

The rest of the paper is organized as follows. In Section 2 we summarize some key results and experimental
predictions. Section 3 outlines the computation of transport coefficients from kinetic theory. Detailed
calculations of transport coefficients at low temperature in any Fermi liquid with an isotropic dispersion
relation are provided in Section 4 in the absence of a background magnetic field, and in Section 5 in
a background magnetic field. We conclude in Section 6 emphasizing our key results and pointing out
experimental implications. Appendices contain a few technical details of our calculations.

2 Main Results

In this paper, we compute the dc thermoelectric transport coefficients of a metal:

$$
\begin{pmatrix}
J_i \\
Q_i
\end{pmatrix} =
\begin{pmatrix}
\sigma_{ij} & T\alpha_{ij} \\
T\alpha_{ij} & T\kappa_{ij}
\end{pmatrix}
\begin{pmatrix}
E_j \\
-\frac{1}{T}\partial_j T
\end{pmatrix},
$$

(3)

where $J_i$ is the charge current, $Q_i$ is the heat current, $E_j$ is an externally applied electric field, and $\partial_j T$
is an “externally applied temperature gradient”. In experiments, one often measures the open circuit
thermal conductivity, where no electrical current flows in the system:

$$
\kappa_{ij} = \bar{\kappa}_{ij} - T\alpha_{ik}\sigma_{kl}^{-1}\alpha_{lj}.
$$

(4)

For much of this paper, we will assume that the conductivity tensors are isotropic: e.g. $\sigma_{ij} = \sigma\delta_{ij}$.

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3 A formal discussion of how this can be done may be found in [5].
A number of systems ranging from graphene \cite{31, 32} to more exotic “strange metals” \cite{10, 33, 34} violate the relations (1) and (2). Some of these violations could arise from the dominance of inelastic scattering processes, while others appear to have a hydrodynamic origin \cite{31}. The purpose of this paper will be to elucidate and expand upon the breakdown of (1) and (2) due to hydrodynamic effects. Indeed, it is well-understood why in the hydrodynamic regime (1) and (2) will fail. For example, a new universal transport relation arises deep in the hydrodynamic regime: \cite{35, 36, 37}

\begin{equation}
\sigma = \frac{e^2 n^2}{T s^2 \kappa} = \frac{-e n}{s} \alpha.
\end{equation}

(5)

Here \( s \) is the entropy density, \( n \) is the number density of electrons, and \(-e\) is the charge of the electron. These ratios emerge because all thermoelectric transport phenomena become linked to momentum-relaxing scattering processes. Also observe that in this hydrodynamic regime, \( \kappa \ll \sigma \) because \( \bar{\kappa}\sigma \approx T\alpha^2 \) \cite{3}; furthermore, \( \bar{\kappa} \gg \kappa \).

The first main result of this paper is the quantitatively accurate description of the transition between the conventional “collisionless” regime (1) and (2), and the hydrodynamic regime (5), carefully accounting for all hydrodynamic effects. In metals with large Fermi surfaces and weak disorder, we will show that

\begin{equation}
\frac{\kappa}{\sigma T} \approx \frac{\pi^2}{3} \frac{\Gamma}{\Gamma + \gamma}.
\end{equation}

(6)

where \( \gamma \) denotes the electron-electron scattering rate, and \( \Gamma \) denotes the electron-impurity scattering rate. While (6) was derived previously in \cite{11}, our derivation will naturally generalize to the Mott law, and to more complicated settings. We will describe the computation of \( \Gamma \) in some detail below, including a qualitative discussion of hydrodynamic effects on \( \Gamma \) \cite{38, 39} in Section 4.5. We briefly discuss finite-temperature corrections to (6) in \( T/T_F \) in Section 4.4. Eq. (6) makes clear that as observed prior, \( \kappa/\sigma T \) is generally smaller than the Wiedemann-Franz prediction in an interacting metal at finite density. Eq. (6) also implies the expected FL behavior at \( T = 0 \), where \( \gamma = 0 \). This is in contrast to a NFL, where it is possible for the WF relation to be violated even at \( T = 0 \). It would be interesting to experimentally study the behavior of the Lorenz number at very low temperatures in strange metals (although in some cases, the putative critical point is unstable to superconductivity, e.g. cuprates, making a low temperature measurement impractical). The recovery of the WF law at low temperatures, or lack thereof, is a simple check on the FL paradigm.

The second main result of this paper is the generalization of (6) to magnetotransport. Including a background magnetic field which leads to cyclotron frequency \( \omega_c \), we find that when \( T \ll T_F \):

\begin{equation}
\frac{\kappa_{xx}}{\sigma_{xx} T} \approx \frac{\pi^2}{3e^2} \left( \frac{\Gamma + \gamma}{\Gamma} \right)^2 + \frac{\omega_c^2}{\Gamma + \gamma^2},
\end{equation}

(7a)

\begin{equation}
\frac{\kappa_{xy}}{\sigma_{xy} T} \approx \frac{\pi^2}{3e^2} \left( \frac{\Gamma^2 + \omega_c^2}{(\Gamma + \gamma)^2 + \omega_c^2} \right).
\end{equation}

(7b)

Like (6), the ratio of the Hall thermal to Hall electrical conductivity strictly decreases with increasing electron-electron interactions. However, the dissipative conductivities exhibit a richer structure. When \( \Gamma + \gamma < \omega_c \), enhancing electron-electron interactions increases the Lorenz number above the ballistic prediction; only when \( \Gamma + \gamma > \omega_c \) does the Lorenz number become smaller than (1). This implies that a non-monotonic temperature dependence of the Lorenz number will occur whenever \( \Gamma < \omega_c \). It is possible, though difficult, to accurately measure electronic thermal conductivity directly \cite{31, 40, 41}. Eq. (7a) is a key prediction for future experiments and will provide a quantitative test of both the hydrodynamic origin of Wiedemann-Franz violations, and the validity of the Fermi liquid paradigm.
In this section, we introduce the general formalism to solve the transport problem. We will start out with rather minimal assumptions beyond the existence of quasiparticles with lifetime $\gg \frac{\hbar}{k_B T}$, and as the section continues we will make more and more specific assumptions. It is (at least in principle) simple to relax many of these assumptions. In practice, going beyond some of these assumptions may necessitate extensive numerical work focusing on specific materials, which is beyond the scope of our current work.

### 3.1 Linearized Boltzmann Equation

In order to compute the thermoelectric conductivity matrix, we solve the Boltzmann equation for a Fermi liquid of dispersion relation $\epsilon(p)$, linearized about thermal equilibrium at temperature $T$. We assume that the band structure is inversion-symmetric and time-reversal invariant. We also assume the presence of an inhomogeneous single-particle potential energy $V_{\text{imp}}(x)$ caused by random impurities. We ignore the spin of the electronic quasiparticles, as we are explicitly interested in spin-independent transport properties, though it is easy to keep track of spin if necessary.

We denote by $f(x,p)$ the distribution function of (quantum) kinetic theory. Letting $\delta f = f_{\text{eq}} + \delta f$, where

$$f_{\text{eq}}(x,p) = \frac{1}{1 + e^{(\epsilon(p) + V_{\text{imp}}(x)-\mu)/k_B T}},$$

denote the distribution function in the absence of any external electric fields or temperature gradients, and $\delta f$ denote the infinitesimal correction arising in infinitesimal electric fields/temperature gradients, we find that the Boltzmann equation reduces to

$$\partial_t \delta f + \mathbf{v} \cdot \nabla \delta f = -\mathbf{F}_{\text{imp}} \cdot \nabla \delta f - \mathbf{F}_{\text{ext}} \cdot \nabla f_{\text{eq}}$$

within linear response. As we are only interested in the dc transport problem, we may drop the time derivative from now on. Here

$$\mathbf{v} = \frac{\partial \epsilon}{\partial p}$$

$\mathbf{F}_{\text{imp}} = -\nabla V_{\text{imp}}$ are the forces arising from the impurity potential, and

$$\delta \mathbf{F}_{\text{ext}} = -e \left( \delta \mathbf{E} - \frac{\epsilon - \mu}{T} \nabla \delta T \right).$$

$\delta C[\delta f]$ denotes the collision integral, written only to leading (first) order in $\delta f$. Explicit discussions of the linearized collision integral may be found in [42]. A solution of the linearized Boltzmann equation (9) will provide $\delta f$. As there are no charge and heat currents in equilibrium, the charge and heat currents are given entirely by this perturbation:

$$J_i(x) = -e \int \frac{d^d p}{(2\pi \hbar)^d} v_i \delta f(x,p),$$

$$Q_i(x) = \int \frac{d^d p}{(2\pi \hbar)^d} (\epsilon + V_{\text{imp}} - \mu) v_i \delta f(x,p),$$

For simplicity in this paper, and without loss of generality, we will shift $\epsilon(p)$ by a constant such that the Fermi surface is located at $\mu = 0$, which simplifies a few of the formulas below.

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[4] Also, the detailed system parameters for carrying out such numerical calculations for specific strongly correlated materials may not be known. For such systems it would not yet be useful to perform serious numerical computations.
Mathematically, the Boltzmann equation is simply a linear algebra problem in a high-dimensional vector space. In order to make analytic progress, we cannot simply invert the linearized Boltzmann operator—it is far too complicated. We instead proceed in a number of steps, which may (for the moment) appear rather cumbersome, but will in the end reduce the calculation of the conductivities to the inversion of few-dimensional matrices, in certain limits. The notation and formulations below follow closely [38, 39].

The first step is to define a suitable inner product for the \( p \)-indices on our vector space:

\[
\langle f | g \rangle = \int \frac{d^d p}{(2\pi \hbar)^d} \left( -\frac{\partial f_{\text{eq}}}{\partial \epsilon} \right) f(p)g(p),
\]

Writing the distribution function as

\[
f(x, p) = f_{\text{eq}}(x, p) + \int \frac{d^d k}{(2\pi)^d} \left( -\frac{\partial f_{\text{eq}}}{\partial \epsilon} \right) \Phi(x, p),
\]

with \( \Phi \) treated as infinitesimally small, and defining

\[
|J_i\rangle = \int d^d p \; v_i(p)|p\rangle,
\]

\[
|Q_i\rangle = \int d^d p \; \epsilon(p) v_i(p)|p\rangle,
\]

we find that

\[
\langle \Phi(x)|J_i\rangle = J_i(x),
\]

\[
\langle \Phi(x)|Q_i\rangle = Q_i(x) - V_{\text{imp}}(x)J_i(x).
\]

The advantage of introducing the inner product (13) is now clear: the highly singular structure of the Fermi function has been absorbed into the integration measure in the inner product. The vectors which we are studying are now smooth functions of momentum across the Fermi surface. We also emphasize that the \( V_{\text{imp}} \)-dependent correction to the heat current will be negligible in the limits studied in this paper. We will think of \(|J_i\rangle\) as the vector that encodes the charge current, and \(|Q_i\rangle\) as the vector that encodes the heat current.

Let \( W \) denote the linearized collision integral. It is a symmetric, positive-semidefinite matrix with the assumptions made above; its null vectors are associated with local conservation laws of electron-electron collisions. We will assume that one of these conservation laws is associated with momentum, because translation invariance is only broken by \( V_{\text{imp}} \). The linearized Boltzmann equation (9) can then be written in our vector notation as

\[
\langle p | \left[ \mathbf{v} \cdot \frac{\partial}{\partial \mathbf{x}} + \mathbf{F}_{\text{imp}} \cdot \frac{\partial}{\partial \mathbf{p}} \right] \Phi \rangle = -\langle p | W |\Phi \rangle + E_i(p|J_i) - \frac{1}{T} \left( \langle p|Q_i \rangle + V_{\text{imp}} \langle p|J_i \rangle \right).
\]

The \((x, p)\)-dependence in the inner product has been properly taken into account—the derivatives above only act on \( \Phi \). Clearly, we may undo the inner product with \(|p\rangle\), since the equation is valid for all \(|p\rangle\).

### 3.2 Perturbative Limit

We now focus on a perturbative limit where the amplitude of the smooth impurity potential \( V_{\text{imp}} \) is perturbatively weak. This is a natural limit of interest for the study of hydrodynamic interaction effects,
where we have defined the matrix

\[
\Phi \approx \begin{pmatrix}
0 & W \\
-1 & 0 \\
\end{pmatrix} + \begin{pmatrix}
i k & 0 \\
W & k & \end{pmatrix} + L_{\text{dis}}(k, k') \begin{pmatrix}
|\Phi(k = 0)\rangle \\
|\Phi(k \neq 0)\rangle \\
\end{pmatrix},
\]

and helps ensure that electron-electron interaction effects are stronger than electron-impurity interactions. Note that although \(V_{\text{imp}}\) is small, it cannot vanish because then strict translational invariance is restored and the electrical conductivity is infinite, in the absence of umklapp (which is ignored throughout this work). This can be seen by writing down Newton’s second law:

\[
\frac{dP_{\text{tot}}}{dt} = Q_{\text{tot}}E
\]

where \(P_{\text{tot}}\) is the total momentum in the theory, and \(Q_{\text{tot}}\) is the total charge. In the absence of translation symmetry breaking, no further terms may be written down on the right hand side, and transport properties in the strict dc limit are ill-posed. In a quantum theory, (18) continues to hold: it is the Ward identity associated with translation symmetry. Therefore, we work consistently in the limit of small, but finite, \(V_{\text{imp}}\).

As is well-known, and we will see explicitly below, all thermoelectric conductivities now scale like \(V_{\text{imp}}^{-2}\) as a consequence of the slowness of momentum relaxation [5, 15, 36, 37]. This means that most – but not all – \(V_{\text{imp}}\)-dependence in the problem can be ignored. To see where we can neglect \(V_{\text{imp}}\), recall that our goal is to compute

\[
J_i^{\text{avg}} = \int \frac{d^d x}{V_d} J_i(x) \approx \int \frac{d^d x}{V_d} \langle \Phi(x)|J_i\rangle
\]

\[
Q_i^{\text{avg}} = \int \frac{d^d x}{V_d} Q_i(x) \approx \int \frac{d^d x}{V_d} \langle \Phi(x)|Q_i\rangle.
\]

Here \(V_d\) denotes the spatial volume – we are simply taking a spatial average. The latter approximations above come from the fact that both \(\sigma\) and \(\kappa\) will scale as \(V_{\text{imp}}^{-2}\) at leading order. It will be useful to Fourier transform \(\Phi\):

\[
|\tilde{\Phi}(k)\rangle \equiv \int \frac{d^d x}{V_d} \Phi(x)e^{-ik\cdot x}.
\]

(9) can be written as

\[
\begin{pmatrix}
W \\
-1 & 0 \\
\end{pmatrix} + \begin{pmatrix}
i k & 0 \\
W & k & \end{pmatrix} + L_{\text{dis}}(k, k') \begin{pmatrix}
|\Phi(k = 0)\rangle \\
|\Phi(k \neq 0)\rangle \\
\end{pmatrix} = \begin{pmatrix} E_i|J_i\rangle - \frac{\partial}{\partial p_i} |Q_i\rangle \end{pmatrix},
\]

where

\[
L_{\text{dis}}(k, k') = -i(k - k')V_{\text{imp}}(k - k')\frac{\partial}{\partial p_i},
\]

and we have denoted \(L_{\text{dis}}^0 = L_{\text{dis}}(0, k)\). Since \(V_{\text{imp}}\) is perturbatively small, and \(ik \cdot v + W\) has no null vectors in general, we do not need to worry about the \(L_{\text{dis}}\) except in the off-diagonal blocks in (21).

Combining all of the above together, we conclude that the thermoelectric conductivity matrix is given by

\[
\begin{pmatrix}
\sigma_{ij} & T\alpha_{ij} \\
T\alpha_{ij} & T\bar{\kappa}_{ij} \\
\end{pmatrix} \approx \begin{pmatrix}
\langle J_i|\bar{W}^{-1}|J_j\rangle & \langle J_i|\bar{W}^{-1}|Q_j\rangle \\
\langle Q_i|\bar{W}^{-1}|J_j\rangle & \langle Q_i|\bar{W}^{-1}|Q_j\rangle \\
\end{pmatrix},
\]

where we have defined the matrix

\[
\bar{W} \approx W + \int \frac{d^d k}{(2\pi)^d} |\langle k_i|k_j\rangle|V_{\text{imp}}(k)|^2 \frac{\epsilon}{\partial p_i} (W + ik \cdot v)^{-1} \frac{\epsilon}{\partial p_j} + O(V_{\text{imp}}^3).
\]

The first term in \(\bar{W}\) corresponds to electron-electron interactions. The second term corresponds to electron-impurity collisions, which will not conserve momentum. Formally speaking, we observe that the
presence of $W$ in the electron-impurity term means that the electron-electron interactions can substantially modify the nature of electron-impurity scattering. This interplay of electron-electron and electron-impurity interactions is described in some detail in [38, 39], and in Section 4.5. However, for most of this paper, we will focus on a limit where $W$ is also taken to be perturbatively small: $W \sim V_{\text{imp}}^2$. In this limit, impurity scattering is well described by the single-particle theory: we may approximate

$$\tilde{W} \approx W + W_{\text{imp}},$$

(25)

where

$$W_{\text{imp}} = \int \frac{d^d k}{(2\pi)^d} k_i k_j |V_{\text{imp}}(k)|^2 \delta(k_i v_i) \frac{\partial}{\partial p_i} \delta(k_j v_j).$$

(26)

Eq. (25) is reminiscent of Matthaisen's rule – the impurity-scattering and electron-scattering diagrams may be added independently to the (disorder-averaged) collision integral of kinetic theory. However, we emphasize that in the final transport coefficients, Mattheisen's rule for resistivities can be violated.

### 3.3 Rotational Invariance and a Basis

The final assumption that we will make in this paper is that the dispersion relation (and disorder distribution) are rotationally invariant. This is a reasonable assumption for many correlated Fermi liquids where the hydrodynamic regime is likely experimentally accessible, including graphene and GaAs. We further assume that $W$ is rotationally invariant; by construction in (26), it is clear that (after performing the $k$-integral) $W_{\text{imp}}$ is rotationally invariant. Thus, $\tilde{W}$ is rotationally invariant. What this means is that the natural basis of vectors $|\mathbf{p}\rangle$ are:

$$|\alpha, m_1 \cdots m_{d-1}\rangle = \int d^d \mathbf{p} f_\alpha(p) Y_{m_1 \cdots m_{d-1}}(\theta_1, \ldots, \theta_{d-1}) |\mathbf{p}\rangle$$

(27)

In the above equation, and henceforth, $p = |\mathbf{p}|$, $\theta_1, \ldots, \theta_{d-1}$ denote the angular coordinates of $\mathbf{p}$, and $f_\alpha$ is a set of polynomial functions that we will detail below. In this basis,

$$\langle \alpha, m_1 \cdots m_{d-1}|\tilde{W}|\beta, m'_1 \cdots m'_{d-1}\rangle = W_{\alpha\beta}^{m_1 \cdots m_{d-1}} \delta_{m_1 m'_1} \times \cdots \times \delta_{m_{d-1} m'_{d-1}}.$$ 

(28)

Our goal is to compute the thermoelectric conductivity matrix. Both $|J_1\rangle$ and $|Q_1\rangle$ are vectors under spatial rotation, and without loss of generality, we may compute $\sigma = \langle J_2|W^{-1}|J_x\rangle$, etc. So a natural choice of basis vectors to study corresponds to

$$|\alpha\rangle \equiv \frac{p_x}{p} f_\alpha(p),$$

(29)

for a basis of polynomials $\{f_\alpha\}$. For these basis vectors, we observe that

$$\langle \alpha|\beta\rangle = \int \frac{d^d \mathbf{p}}{(2\pi \hbar)^d} f_\alpha(p) f_\beta(p) \frac{p_x^2}{p^2} = \frac{1}{d} \int \frac{d^d \mathbf{p}}{(2\pi \hbar)^d} f_\alpha(p) f_\beta(p).$$

(30)

Up to the overall prefactor of $1/d$, we may simply compute the radial integrals over the Fermi surface.

Another useful simplification in this basis arises when computing the matrix elements $\langle \alpha|W_{\text{imp}}|\beta\rangle$. Using

$$k_i \frac{\partial}{\partial p_i} \left( \frac{p_x}{p} f_\alpha \right) = \frac{k_x}{p} f_\alpha + (f_\alpha - 1) \frac{k_i p_i p_x}{p^2},$$

(31)

and observing that the $\delta(k_i v_i)$ in (26) constrains $k_i p_i = 0$ in an isotropic theory, we conclude that

$$\langle \alpha|W_{\text{imp}}|\beta\rangle = \Omega_d \int \frac{p^{d-1} dp}{(2\pi \hbar)^d} \left( - \frac{\partial f_{\text{eq}}}{\partial \varepsilon} \right) \int \frac{d^d \mathbf{k}}{(2\pi \hbar)^d} k_x^2 |V_{\text{imp}}(\mathbf{k})|^2 \delta(k_i v_i) \frac{f_\alpha f_\beta}{p^2}.$$
\[ \Omega_d \int \frac{p^{d-1} dp}{(2\pi \hbar)^d} \left( -\frac{\partial f_{eq}}{\partial \epsilon} \right) f_\alpha f_\beta \frac{1}{p^2 v} \left[ \int \frac{d^d k}{(2\pi)^d} k^2 |V_{\text{imp}}(k)|^2 \delta(k_x) \right] \]

\[ = A_{\text{imp}} \times \Omega_d \int \frac{p^{d-1} dp}{(2\pi \hbar)^d} \left( -\frac{\partial f_{eq}}{\partial \epsilon} \right) f_\alpha f_\beta \frac{1}{p^2 v}, \]

where \( \Omega_d \) is the area of the unit sphere. Hence, all of the physics of impurity scattering becomes captured by a single constant \( A_{\text{imp}} \), which is defined as the object in square brackets in the second line of (32). We caution that there can be non-trivial temperature dependence in \( A_{\text{imp}} \), in the presence of temperature-dependent screening of a long-range Coulomb impurity potential \( V_{\text{imp}}(k) \) [43, 44], an effect that we ignore in the present work.

## 4 Large Fermi Surfaces

To make further progress, we will now work in the limit where the Fermi surface is large. To be precise, we will assume that \( T \ll p_F v_F \). In this limit, we can further simplify the evaluation of the conductivity by neglecting all but a few low order polynomials \( f_\alpha \) in the basis (29).

### 4.1 An Efficient Basis

Let the dispersion relation be

\[ \epsilon(p) = v_F q + \frac{a}{2} q^2 + \frac{b}{3} q^3 + \cdots, \]  

(33)

A natural set of basis functions can be constructed as follows. Letting \( q \equiv p - p_F \), we define the states \( |q^n\rangle \), corresponding to the polynomials \( f_\alpha = q^n \). Of great importance to us is that in this basis,

\[ |J_x\rangle = -e \left( v_F |q^0\rangle + a |q^1\rangle + b |q^2\rangle + \cdots \right), \]  

(34a)

\[ |Q_x\rangle = v_F^2 |q^1\rangle + \frac{3}{2} v_F a |q^2\rangle + \cdots. \]  

(34b)

Unfortunately, the basis functions \( |q^n\rangle \) are not normalized. It would be more convenient if we could work with an orthonormal basis. This can be constructed straightforwardly by the Gram-Schmidt procedure, and we will denote the result of this procedure with \( |n\rangle \). The calculation of \( |n\rangle \) from \( |q^n\rangle \) is straightforward but tedious: details are provided in Appendix A. A useful qualitative fact for the discussion that follows is that

\[ \langle q^n | q^m \rangle \sim \begin{cases} T^{n+m} & \text{n + m even} \\ T^{n+m+1} & \text{n + m odd} \end{cases}. \]  

(35)

In what follows, subleading numerical coefficients in \( T \) are specific to \( d = 2 \) spatial dimensions, but the general structure is unchanged for all dimensions. The first few basis functions are

\[ |0\rangle = c_{00} \sqrt{\frac{d}{\nu}} |q^0\rangle, \]  

(36a)

\[ |1\rangle = \sqrt{\frac{d}{\nu}} \left[ \frac{c_{11}}{T} |q^1\rangle + T c_{10} |q^0\rangle \right], \]  

(36b)

with \( \nu \) the density of states (we remind the reader of useful thermodynamic formulas for a Fermi liquid in the low temperature limit in Appendix B), the factor of \( d \) arising from (30), and

\[ c_{00} = 1 - \frac{\pi^2 T^2}{2} \left( \frac{a^2}{2 v_F^4} - \frac{a}{2 p_F v_F^3} - \frac{b}{3 v_F^3} \right) + O(T^4), \]  

(37a)
\[
\begin{align*}
\mathcal{C}_{11} &= \frac{\sqrt{3}v_F}{\pi} + \frac{\pi T^2 (2v_F^2 + 14bv_Fp_F^2 + 15ap_Fv_F - 27a^2p_F^2)}{4\sqrt{3}v_F^3p_F^2} + O(T^4), \\
\mathcal{C}_{10} &= \frac{\pi (3ap_F - 2v_F)}{2\sqrt{3}p_Fv_F^2} + O(T^2).
\end{align*}
\] (37b)

A general basis function is
\[
\langle n \rangle = \sqrt{\frac{d}{\nu}} \left( \frac{v_F}{T} \right)^n c_{nn}\langle q^n \rangle + \left( \frac{v_F}{T} \right)^{n-2} (c_{n,n-1}\langle q^{n-1} \rangle + c_{n,n-2}\langle q^{n-2} \rangle) + \cdots .
\] (38)

A key observation is that – up to \(O(T^2/T_F^2)\) corrections – the orthonormal basis is equivalent to \(\langle q^n \rangle\), up to a prefactor which is proportional to \(T^{-n}\). Also observe that \(\langle q^n \rangle\) can exactly be expressed as a function of \(\langle m \rangle\) with \(m \leq n\).

### 4.2 Approximating the Collision Integrals

From the form of (34) and (36), it is clear that as long as components of \(W_{\text{imp}}\) are “well-behaved” in the low \(T\) limit, then we can efficiently compute the thermoelectric conductivity matrix by focusing only on the first few basis vectors: \(\langle 0 \rangle, \langle 1 \rangle, \ldots \). Indeed, this is the case:
\[
\langle m | W_{\text{imp}} | n \rangle \lesssim T^{\text{m-n}}.
\] (39)

To show this, we first write
\[
\frac{A_{\text{imp}}}{p^2v(p)} = \Gamma_1 q + \Gamma_2 q^2 + \cdots .
\] (40)

By definition, \(q^k\langle q^n \rangle = \langle q^{n+k} \rangle\). (We are denoting the linear operation which multiplies by \(q\) as \(q\), in the obvious way.) Now, assuming that \(m > n\), without loss of generality, observe that
\[
\langle m | q^{m-n} | n \rangle \sim \langle m | \left[ \frac{1}{T^n}\langle q^m \rangle + \frac{T^2}{T^n}\langle q^{m-1} \rangle + \frac{T^2}{T^n}\langle q^{m-2} \rangle + \cdots \right] \sim \langle m | q^m \rangle \sim T^{m-n}.
\] (41)

In fact, staring at the above expression, we also find that \(\langle m - \ell | q^{m-n} | q^n \rangle \lesssim T^{m-n} \) for any \(\ell \geq 0\). Therefore,
\[
\langle m | W_{\text{imp}} | n \rangle \sim \Gamma_{m-n} T^{m-n} + \cdots .
\] (42)

The \(\cdots\) in (42) includes terms proportional to \(\Gamma_{m-n+2}T^{m-n+2}\), etc., which are all subleading in the low temperature limit. At \(T = 0\), there is a residual resistivity:
\[
\langle m | W_{\text{imp}} | n \rangle = \Gamma \delta_{mn}.
\] (43)

We have not dropped any constant prefactor in the above formula (43).

We next observe that
\[
\langle m | W_{\text{imp}}^{-1} | n \rangle \lesssim T^{\text{m-n}}.
\] (44)

One way to show this result is recursively. From the form of the \(T = 0\) result (43), we show the base case \(|n - m| = 0\). For finite \(|n - m|\), we may use block matrix inversion identities. Without loss of generality, we again take \(m > n\). Let us write
\[
W = \left( \begin{array}{cc} \langle k < m | & \langle k \geq m | \end{array} \right) \left( \begin{array}{cc} W_{--} & W_{+-} \\ W_{+} & W_{++} \end{array} \right) \left( \begin{array}{c} |k < m \rangle \\ |k \geq m \rangle \end{array} \right)
\] (45)
Since there are no singular eigenvalues at low temperatures,
\[
\langle m | W_{\text{imp}}^{-1} | n \rangle \sim \sum_{k<n, \ell \geq m} (W_{++} - W_{+-} W_{-+})^{-1}_{m\ell} (W_{+-})_{k\ell} (W_{-+})_{kn} \lesssim \sum_{k<n, \ell \geq m} T^{k-\ell + |k-n|} \lesssim T^{m-n}.
\]

Since physical observables such as \(|J_x\) and \(|Q_x\) are naturally expressed in terms of the \(|q^n\) basis, as in (34), we are only interested in evaluating \(W_{\text{imp}}^{-1}\) on vectors which take the form
\[
|J_x\rangle = \sum_n a_n T^n |n\rangle, \quad |Q_x\rangle = \sum_n b_n T^n |n\rangle, \quad \text{etc.}
\]
Observing that
\[
\sum m a_m T^{m+n} \langle m | W_{\text{imp}}^{-1} | n \rangle \sim \sum m a_m a_n T^{m+n+|m-n|} = \sum m a_m a_n T^{2 \max(m,n)},
\]
we conclude that to compute a transport coefficient up to \(O(T^2)\), we only need to keep basis vectors \(|0\rangle, \ldots, |k\rangle\). To leading order, we know from the Wiedemann-Franz law and Mott law that \(T\bar{\kappa}\) and \(T\alpha\) will both scale as \(T^2\) at low temperatures. As we show in Section 4.3, keeping only the \(|0\rangle\) and \(|1\rangle\) basis vectors is sufficient to recover these relations in the non-interacting limit.

Now that we understand the form of the “collisionless” electron-impurity scattering, let us discuss the form of the electronic collision integral. Since we are interested in the hydrodynamic regime of electron flow, we focus on theories where momentum is conserved in all electron-electron collisions (there is no umklapp). This means that the momentum vector
\[
|P_x\rangle = p_F |q^0\rangle + |q^1\rangle \approx \sqrt{\nu \frac{\pi T}{d}} \left[ p_F |0\rangle + \frac{\pi T}{\sqrt{3}} |1\rangle \right]
\]
is a null vector of \(W\). One simple choice of \(W\) that is consistent with this requirement is
\[
W \approx \gamma \left( |1\rangle - \frac{\pi T}{\sqrt{3} p_F v_F} |0\rangle \right) \left( |1\rangle - \frac{\pi T}{\sqrt{3} p_F v_F} |0\rangle \right) + \sum_{n=2}^\infty \gamma |n\rangle \langle n|.
\]
This choice, which is used extensively in transport theories, is often called the relaxation time approximation [45]. Note that the relative decay of longitudinal fluctuations above necessitates inelastic scattering processes. The coefficient \(\gamma \approx \tau^{-1}_{ee}\) is associated with the rate at which electron-electron collisions relax longitudinal fluctuations of the distribution function, proportional to \(p_x\):
\[
\gamma \equiv \tilde{\alpha}^2 \frac{T^2}{T_F}.
\]
where \(\tilde{\alpha}^2\), an effective coupling constant in the problem which we do not specify, is (roughly) proportional to the interaction constant. In lower dimensions, it is possible for this scattering time to be enhanced by factors of \(\log(T_F/T)\) [46, 47, 48, 49, 50, 51], and we will not worry about these logarithms in this paper.

We note that the Fermi liquid prediction (51) for the inelastic scattering rate – with suitable logarithms – has been verified experimentally (see e.g. [52] for a direct verification in a 2D GaAs system). The \(T^2\) dependence in (51) is completely consistent with the classic Fermi liquid paradigm of well-defined quasiparticles. Indeed, our kinetic picture is perturbative in interactions and so a NFL does not emerge in our model in 2D or 3D electron liquids. This statement is independent of \(\tilde{\alpha}^2\), although for large enough \(\tilde{\alpha}^2\) our kinetic theory is not valid. Normal 3D metals are strongly interacting in the sense that \(\tilde{\alpha}\) (also commonly called ‘\(\tau_{fs}\)’ in interacting Coulomb systems) \(\sim 6\). However, because \(T_F \sim 10^4\) K, \(\gamma\) is still small relative to \(k_B T/h\) (the scattering rate below which a quasiparticle picture does not make sense [5]). In 2D GaAs hole systems, the interaction coupling constant can be \(\sim 40\) in dilute experimental systems, putting the system deep in the hydrodynamic regime [53].
4.3 The Strongly Interacting Limit

We are now finally ready to study transport phenomena across the ballistic-to-hydrodynamic crossover. To start off, let us work in a simple limit where

\[ \Gamma \ll \alpha^2 T_F. \]  

(52)

We will relax this assumption in Section 4.4. This inequality can be satisfied in 2D systems experimentally, but not in the usual metals. (52) implies that \( \gamma = \Gamma \) at temperatures \( T \ll T_F \), where an expansion in low order basis vectors ought to be sensible, even in the hydrodynamic regime. Thus, we will work only to the lowest non-trivial order in \( T/T_F \) in \( \mathcal{W}_{\text{imp}} \), to capture the effects of interactions. From the form of the Wiedemann-Franz law (1), Mott relation (2), and the definition (23), we conclude that it will suffice to keep track of only the vectors \( |0\rangle \) and \( |1\rangle \). This is not equivalent to a Taylor series expansion in \( T \), as \( \gamma \) itself depends on \( T \) via (51). We will return to this point in Section 4.4.

Converting the \( |q^n\rangle \) basis to the \( |n\rangle \) basis, the charge and heat current vectors are

\[ |J_x\rangle \approx -e \sqrt{\nu d} \left[ \frac{v_F}{\sqrt{3}} |0\rangle + \frac{\pi T v_F}{\sqrt{3}} |1\rangle \right], \]  

(53a)

\[ |Q_x\rangle \approx \sqrt{\nu d} \left[ \frac{\pi^2 T^2}{3 p_F^2} |0\rangle + \frac{\pi T v_F}{\sqrt{3}} |1\rangle \right]. \]  

(53b)

The net collision integral \( \tilde{W} \) is given by

\[ \tilde{W} \approx \left( \frac{\pi^2 T^2}{3 p_F^2 v_F^2} \gamma + \Gamma \right) |0\rangle\langle 0| + \frac{\pi T}{\sqrt{3} v_F} \left( \frac{\Gamma_1 - \gamma}{p_F} \right) (|0\rangle\langle 1| + |1\rangle\langle 0|) + \frac{\pi^2 T^2}{3 p_F^2 v_F^2} \gamma + \frac{\pi^2 T^2}{3 p_F^2 v_F^2} \gamma + \frac{\pi T}{\sqrt{3} v_F} \left( \frac{\Gamma_1 - \gamma}{p_F} \right) (|0\rangle\langle 1| + |1\rangle\langle 0|) \]  

(54)

Computing the thermoelectric conductivity matrices has reduced to simply inverting a \( 2 \times 2 \) matrix:

\[ \tilde{W}^{-1} \approx \frac{1}{\Gamma} |0\rangle\langle 0| - \frac{\pi T}{\sqrt{3} v_F} \left( \frac{\Gamma_1 - \gamma}{p_F} \right) (|0\rangle\langle 1| + |1\rangle\langle 0|) + \frac{\pi^2 T^2}{3 p_F^2 v_F^2} \gamma + \frac{\pi^2 T^2}{3 p_F^2 v_F^2} \gamma + \frac{\pi T}{\sqrt{3} v_F} \left( \frac{\Gamma_1 - \gamma}{p_F} \right) (|0\rangle\langle 1| + |1\rangle\langle 0|) \]  

(55)

In this step, we have made approximations consistent with the assumptions stated at the start of this subsection.

First, let us study the limit \( \gamma = 0 \), which corresponds to the \( T \to 0 \) limit. After some algebra we obtain

\[ \sigma \approx \frac{\nu e^2 v_F^2}{d \Gamma}, \]  

(56a)

\[ \kappa \approx \tilde{\kappa} \approx \frac{\pi^2 T \nu v_F^2}{3 d \Gamma} \equiv \kappa_{\text{non-int}}, \]  

(56b)

\[ \alpha \approx \frac{\pi^2 T \nu}{3 d} \left[ -v_F \frac{\Gamma_1}{\Gamma^2} + \frac{a}{\Gamma} + \frac{v_F}{p_F \Gamma} \right] \equiv \alpha_{\text{non-int}}. \]  

(56c)

In each case, we have only written down the leading order terms in the \( T \to 0 \) limit (where, by definition, we should recover the standard Fermi liquid WF law since interaction-induced scattering vanishes at the Fermi surface). It is clear that we recover the Wiedemann-Franz law (1). The Mott relation (2) is also obeyed, because

\[ \frac{\partial \sigma}{\partial \mu} = \frac{1}{d v_F} \frac{\partial \sigma}{\partial p_F} = \frac{\Omega_d}{(2\pi)^4 d v_F} \frac{\partial}{\partial p_F} \frac{p_F v_F}{\Gamma} = \frac{\Omega_d}{(2\pi)^4 d v_F} \left[ \frac{v_F}{\Gamma} - \frac{p_F v_F}{\Gamma^2} \frac{p_F a}{\Gamma} \right] \]
\[ \text{(57)} \]

Having confirmed that our theory correctly reproduces the non-interacting “collisionless” limit of electron-impurity scattering, let us crank up the electron-electron collision rate \( \gamma \). The first thing we observe is that

\[
\sigma \approx \nu e^2 v_F^2 \Gamma + O(T^2, T^2 \gamma). \tag{58}
\]

This is an important result: interactions affect the electrical conductivity only weakly (i.e. at higher-order terms) through hydrodynamic corrections. By contrast, as we show below, interaction does affect the thermal conductivity and the thermoelectric conductivity. The electrical conductivity is, in this approximation, essentially independent of electron-electron scattering.

Next, let us discuss the thermal conductivity \( \bar{\kappa} \). In this case, we obtain

\[
\bar{\kappa} = \nu \pi^2 v_F^2 T \left[ \frac{\pi^2 T^2}{3 v_F^2} \gamma + \Gamma - 2 \frac{\pi^2 T^2}{3 v_F^2} \left( \frac{\Gamma}{\Gamma + \gamma} \right) \Gamma + \gamma \right] \approx \kappa_{\text{non-int}} \Gamma \frac{\Gamma}{\Gamma + \gamma} + \kappa_{\text{int}} \Gamma \frac{\Gamma}{\Gamma + \gamma}, \tag{59}
\]

with

\[
\kappa_{\text{int}} = \frac{4\pi^2 T^3 \nu}{9 d v_F^2}. \tag{60}
\]

The thermoelectric conductivity is given by

\[
\alpha \approx -e \frac{\nu \pi^2 T^2 v_F}{3d} \left[ \frac{\pi^2 T^2}{3 v_F^2} \gamma + \Gamma - \frac{\pi^2 T^2}{3 v_F^2} \left( \frac{\Gamma}{\Gamma + \gamma} \right) \Gamma + \gamma \right] \approx \alpha_{\text{non-int}} \Gamma \frac{\Gamma}{\Gamma + \gamma} + \alpha_{\text{int}} \Gamma \frac{\Gamma}{\Gamma + \gamma}, \tag{61}
\]

where

\[
\alpha_{\text{int}} = -e \frac{2\pi^2 T \nu v_F}{3d v_F^2}. \tag{62}
\]

Both \( \bar{\kappa} \) and \( \alpha \) admit an elegant interpretation in this limit: they are a weighted average of the “collisionless” transport coefficient and an “interaction-limited” transport coefficient. The weights in front of each term correspond to the fraction of the total scattering rate that is in the electron-impurity vs. electron-electron channel.

We can go further. In the interaction-dominated limit, we expect that the thermoelectric conductivity matrix is limited entirely by momentum relaxation [5, 15, 35, 36, 37]. Writing down the momentum conservation equation, we obtain

\[
-e n \mathbf{E} - s \nabla T = \Gamma_\ast \mathbf{v}, \tag{63}
\]

with \( \mathbf{v} \) the velocity of the interacting fluid, and \( \Gamma_\ast \) a coefficient related to weak momentum relaxation. The left hand side corresponds to the momentum added to the system by the external drives; the right hand side corresponds to the system’s internal momentum dissipation. Approximating the charge current as \( \mathbf{J} \approx n \mathbf{v} \) and the heat current as \( \mathbf{Q} \approx T s \mathbf{v} \), we arrive at the formulas

\[
\left( \begin{array}{cc} \sigma & T \alpha \\ T \alpha & T \bar{\kappa} \end{array} \right) \approx \frac{1}{\Gamma_\ast} \left( \begin{array}{cc} e^2 n^2 & -e T s n \\ -e T s n & (T s)^2 \end{array} \right). \tag{64}
\]

Using the thermodynamic relations of a low temperature Fermi liquid given in Appendix B, we find that in our toy model above,

\[
\Gamma_\ast = M \Gamma, \tag{65}
\]

where \( M \) is the momentum-momentum susceptibility (analogous to a mass density). Since \( \Gamma \) is the momentum relaxation rate, this agrees with the predictions of the memory matrix formalism [36, 37].
Using (60) and (62) with these thermodynamic identities, we can confirm that \( \alpha_{\text{int}} \) and \( \bar{\kappa}_{\text{int}} \) take the form demanded by (64), relative to \( \sigma \). Note also the identity

\[
\sigma \bar{\kappa}_{\text{int}} = T \alpha_{\text{int}}^2, \tag{66}
\]

which is tied to the fact that one process – momentum relaxation – limits all transport coefficients in the hydrodynamic regime.

Now, let us return to the fate of the Wiedemann-Franz law. Conventionally one compares not \( \bar{\kappa} \) to \( \sigma \), but \( \kappa \) to \( \sigma \). As a consequence, using the identity (66), we obtain

\[
\kappa \approx \frac{\Gamma}{\Gamma + \gamma} \kappa_{\text{non-int}} + \frac{\gamma \Gamma}{(\Gamma + \gamma)^2} \left( \bar{\kappa}_{\text{int}} - \frac{2T \alpha_{\text{int}} \alpha_{\text{non-int}}}{\sigma} \right) \approx \frac{\Gamma}{\Gamma + \gamma} \kappa_{\text{non-int}} \approx \frac{\pi^2 T \nu v_F^2}{3d(\Gamma + \gamma)} \tag{67}
\]

As before, we are neglecting contributions that are subleading in \( T \), at each order in \( \gamma \). The Wiedemann-Franz law will be violated, “strongly interacting” limit, in the particularly elegant manner given in (6). In this simple model, the amount of violation of the Wiedemann-Franz law corresponds directly to the fraction of scattering processes which are momentum-conserving. This interpretation fails to hold if \( T \) is not so small, and/or if the disorder strength is comparable to the interaction strength (Section 4.4). Earlier discussions emphasizing that the Wiedemann-Franz law will be violated in the hydrodynamic limit of a finite density metal, because \( \kappa/\sigma T \) is very small, can be found in [3, 11, 54]. In the hydrodynamic limit, the value of \( \kappa \) matches the dissipative hydrodynamic coefficient \( \kappa_\alpha \), at leading order in \( T/T_F \): see Appendix D.

The fate of the Mott relation is less dramatic. In the limit \( T \ll T_F \), both \( \alpha_{\text{non-int}} \) and \( \alpha_{\text{int}} \) are linear in \( T \). The collisionless-to-hydrodynamic crossover will simply correspond to a gradual change in the slope of \( \alpha(T) \). We note that this crossover can be more dramatic near charge neutrality (\( T \gtrsim T_F \)). Some experimental evidence for this effect in graphene, and further discussion, can be found in [32, 55].

### 4.4 The Weakly Interacting Limit

In this section, we will consider the limit where (52) is not obeyed. This corresponds to metals with relatively low Fermi temperature (where the WF law is affected by thermal smearing of the Fermi function even without any interaction effects), and/or relatively small interaction coupling \( \bar{\alpha} \). Indeed, smearing of the Fermi surface leads to comparable \( O(T^2) \) corrections, similar to (but distinct from) electron-electron interactions.

To correctly compute the thermal conductivity, we must compute \( T \bar{\kappa} \) to \( O(T^4) \), and so we must keep the basis vector \( |2 \rangle \) in our expansion. For simplicity, we will take the dispersion relation \( \epsilon(p) \) to be exactly quadratic. This means that in (33), \( b = 0 \), \( a p_F = v_F \), and \( p_F v_F = 2T_F \). We will also treat electron-electron collisions within a relaxation time approximation (50), but at next-to-leading order this does not change the form of the answer. The forms of \( |J_x \rangle \), \( |Q_x \rangle \) and \( W_{\text{imp}} \) are written explicitly in Appendix C. We then explicitly compute

\[
\sigma = \frac{\nu v_F^2}{2} \left( 1 + \frac{\pi^2 T^2}{3 T_F^2} \right) + O(T^4), \tag{68a}
\]

\[
\kappa = \frac{\nu \pi^2 T v_F^2}{2} \left( 1 - \frac{17 \pi^2 T^2}{30 T_F^2} - \frac{\bar{\alpha}^2 T^2}{T T_F} \right) + O(T^5). \tag{68b}
\]

The Lorenz number

\[
\frac{\kappa}{\sigma T} = \frac{\pi^2}{3 e^2} \left[ 1 - \frac{9 \pi^2 T^2}{10 T_F^2} - \frac{\bar{\alpha}^2 T^2}{T T_F} \right] + \cdots \tag{69}
\]

is thus decreased both by thermal smearing of the Fermi surface and by electron-electron interactions. When disorder is very weak, the dominant effect will be that of electron-electron interactions. Also observe
that even the first subleading temperature-dependent correction to $\sigma$ arises only from thermal smearing of the Fermi surface, and not from electron-electron scattering.

Thus, $\Gamma$, $\gamma$, and $T_F$ are all relevant parameters determining the effective Lorenz number as a function of $T$. Remembering that $T_F \sim n^{2/d}$, where $n$ is the electron density of the $d$-dimensional electron liquid, and that $\gamma \sim T^2/T_F$, we conclude that lowering electron density may enhance both the effect of interaction and the effect of Fermi surface thermal smearing depending on the temperature of the system. If the effective impurity disorder Gamma itself has a temperature dependence through electronic screening (as it does in 2D systems [56]), the situation gets much more complicated. Disentangling all of these effects quantitatively will require materials-specific computations beyond the scope of the present paper.

4.5 Hydrodynamic Fluctuations and Impurities

In this section, we will qualitatively describe the interplay of electron-electron interactions with the impurity potential. Mathematically, this is done by keeping track of the full $W$ matrix when performing the matrix inverse in (24). For the purposes of this section, we will keep the discussion qualitative, since the quantitative details depend on far too many unknown material-specific parameters and the precise nature of the disorder [38, 39]. After a microscopic calculation, what one finds is that $W_{\text{imp}}$ is no longer approximately a single scattering rate $\Gamma$, times the identity matrix. Instead, the eigenvalues of $W_{\text{imp}}$ will sensitively depend on the nature of disorder, and on the modes $|\alpha\rangle$ which are being sourced.

We assume that the inhomogeneous potential $V_{\text{imp}}(x)$ varies on the length scale $\xi$. Let $\theta \ll 1$ denote the typical deflection angle of a quasiparticle after it moves a distance $\sim \xi$ through the potential landscape. A typical degree of freedom $|\alpha\rangle$ will obey

$$\Gamma \sim \langle \alpha | W_{\text{imp}} | \alpha \rangle \sim \theta^2 \frac{v_F \ell_{\text{ee}}}{\xi (\xi + \ell_{\text{ee}})},$$

where

$$\ell_{\text{ee}} \equiv \frac{v_F}{\gamma}.$$  

This will hold, at least qualitatively, for all modes $|0\rangle$, $|1\rangle$, $|2\rangle$, etc., in the isotropic Fermi liquid with potential disorder [38]. In these theories, $\Gamma$ decreases as electron-electron interactions increase [38]:

$$\Gamma \approx \theta^2 \frac{v_F}{\xi} \left[ 1 - \frac{\xi \gamma}{v_F} + \ldots \right] \approx \theta^2 \frac{v_F}{\xi} - \theta^2 \gamma + O(\gamma^2),$$

We conclude from (6) that the Wiedemann-Franz law is violated even more strongly when hydrodynamic corrections to impurity scattering are accounted for, since the effective disorder strength $\Gamma$ is suppressed by interactions. From (72), we also estimate that the hydrodynamic corrections to impurity scattering are relatively weak compared to intrinsic electron-electron scattering in the collision integral by a factor $\theta^2$. This arises from the inherent difference between elastic (impurity) and inelastic (electron-electron) scattering processes.

However, in some cases it is possible for hydrodynamic corrections to impurity scattering to behave rather differently. In particular, in the presence of “magnetic” disorder, one obtains [39]

$$\langle 0 | W_{\text{imp}} | 0 \rangle \approx \theta^2 v_F \left( \frac{1}{\xi} + \frac{1}{\ell_{\text{ee}}} \right) = \theta^2 \frac{v_F}{\xi} + \theta^2 \gamma \equiv \Gamma_*.$$  

$|1\rangle$, $|2\rangle$, etc. will have the same decay rate as before. The origin of the peculiar behavior of the decay rate of the $|0\rangle$ mode is related to the fact that the “magnetic” disorder exerts large shear stresses which

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5This relation changes for systems with non-parabolic dispersion relations, such as monolayer graphene.
in turn create large local velocity fields in the electronic fluid. We must now recompute $\sigma$ and $\kappa$ in a fluid where
\[ W_{\text{imp}} = \Gamma_* |0\rangle \langle 0| + \Gamma |1\rangle \langle 1|. \tag{74} \]
Following the procedure of Section 4.3, we obtain
\[
\sigma \approx \frac{\nu e^2 v_F^2}{d \Gamma_*}, \tag{75a}
\]
\[
\kappa \approx \frac{\nu \pi^2 T}{3} \frac{v_F^2}{\Gamma + \gamma}. \tag{75b}
\]
We therefore find that for this fluid,
\[
\theta^2 \lesssim \frac{3e^2}{\pi^2} \frac{\kappa}{\sigma T} \leq 1. \tag{76}
\]
The Wiedemann-Franz law is still violated, but not as strongly as before. Furthermore, the Lorenz ratio will no longer become arbitrarily small as the sample becomes more pure. This scenario is analogous to the presence of electron-electron umklapp (momentum-relaxing) scattering at rate $\theta^2 \gamma$.

Finally, another well-known [56] example of an interaction-suppressing effect of disorder is the screening of long range Coulomb impurities by the electrons themselves, which leads to a strongly temperature-dependent effective disorder in 2D systems, manifesting a first-order positive thermal correction to the resistivity which is linear in $T/T_F$ in 2D (but not in 3D). This effect, which is beyond the scope of the current hydrodynamic theory as it arises from the singular nature of the 2D polarizability function at $2k_F$ [57, 58], tends to increase $\Gamma$, and therefore the Lorenz number, with increasing temperature. As this effect will increase the Lorenz number it does not provide an alternate explanation for reduced Lorenz numbers in an interaction-limited regime.

5 Magnetotransport

In this section, we will discuss magnetotransport phenomena in the presence of a weak magnetic field, focusing specifically on the experimentally relevant case of a two-dimensional electron fluid. When the cyclotron radius is very large compared to $\lambda_F$, the magnetic field can be treated within our classical kinetic description. If the cyclotron radius is large compared to the size of the impurity potential, then we can neglect magnetic field corrections to $W_{\text{imp}}$. We will make both of these assumptions to simplify the calculations below.

Accounting for the magnetic field amounts to modifying $\tilde{W}$ to $\tilde{W} = W + W_{\text{imp}} + W_{\text{mag}}$, with
\[
W_{\text{mag}} = -eB \epsilon_{ij} v_i \frac{\partial}{\partial p_j}, \tag{77}
\]
and $\epsilon_{xy} = -\epsilon_{yx} = 1$ the antisymmetric Levi-Civita tensor. In a rotationally invariant Fermi liquid, we can further simplify this. Generalizing (29) to
\[
|\alpha_i\rangle \equiv \frac{p_i}{p} f_\alpha(p), \tag{78}
\]
and generalizing the bases discussed in Section 4.1, we obtain that
\[
W_{\text{mag}}|\alpha_i\rangle = e\epsilon_{ij} \frac{p_j}{p} \frac{Bv(p)}{p} f_\alpha(p); \tag{79}
\]
hence, $W_{\text{mag}}$ will mix $x$ and $y$ vectors (under spatial rotation), but will not couple vectors to higher/lower rank tensors.
5.1 The Strongly Interacting Limit

For simplicity, let us now focus on the strongly interacting limit described in Section 4.3. As before, we will (up to the \( T \)-dependence of \( \gamma \)) only be interested in the leading order temperature dependence of all conductivities, which will allow us to neglect all basis vectors but \(|0_{x,y}\rangle\) and \(|1_{x,y}\rangle\). We may write

\[
\tilde{\mathcal{W}} = (\mathcal{W} + \mathcal{W}_{\text{imp}}) \otimes \delta_{ij} + \tilde{\mathcal{W}}_{\text{mag}} \otimes \epsilon_{ij},
\]

where

\[
\tilde{\mathcal{W}}_{\text{mag}} = \frac{eBv_F}{p_F} (|0\rangle\langle 0| + |1\rangle\langle 1|) + \frac{\pi T eB(ap_F - v_F)}{\sqrt{3}v_F p_F^2} (|0\rangle\langle 1| + |1\rangle\langle 0|).
\]

Using (23), we can invert the \( 4 \times 4 \) matrix \( \tilde{\mathcal{W}} \) to compute all conductivities of interest. Taking the same limit as Section 4.3, we find that the electrical conductivity is given by

\[
\sigma_{xx} = \sigma_{yy} = \nu \frac{e^2 v_F^2 \Gamma}{d (\Gamma^2 + \omega_c^2)},
\]

\[
\sigma_{xy} = -\sigma_{yx} = \nu \frac{e^2 v_F^2 \omega_c}{d (\Gamma^2 + \omega_c^2)},
\]

where we have defined the cyclotron frequency

\[
\omega_c = \frac{eBv_F}{p_F}.
\]

As before, we observe that \( \sigma_{ij} \) is independent of electron-electron interactions, within this simple model. Furthermore, the form of \( \sigma_{ij} \) is completely consistent with the Drude model of magnetotransport: see e.g. [15]. The resistivity tensor \( \rho_{ij} = \sigma_{ij}^{-1} \) exhibits no classical magnetoresistance, following the conventional lore. When \( \Gamma = 0 \), we obtain the Hall conductivity

\[
\sigma_{xy} = \sigma_{yx} = \frac{\nu e^2 v_F^2}{2 \omega_c} = \frac{en}{B}.
\]

Since \( -en \) is the charge density, we find the classical Hall conductivity. In fact, using quantum Ward identities, this relation can be derived for any translation invariant quantum system (in the absence of Berry curvature): see e.g. [5]. The open-circuit thermal conductivities are given by

\[
\kappa_{xx} = \kappa_{yy} = \frac{\nu \pi^2 T v_F^2 (\Gamma + \gamma)}{d 3 (\Gamma^2 + \omega_c^2 + \omega_c^2)},
\]

\[
\kappa_{xy} = -\kappa_{yx} = -\frac{\nu \pi^2 T v_F^2 \omega_c}{d 3 (\Gamma^2 + \omega_c^2 + \omega_c^2)},
\]

while the closed-circuit thermal conductivity is

\[
\bar{\kappa}_{xx} = \bar{\kappa}_{yy} \approx \kappa_{xx} + \frac{4\pi^4 T^3 \Gamma^2 \gamma^2}{d p_F^2 (\Gamma^2 + \omega_c^2)((\Gamma + \gamma)^2 + \omega_c^2)},
\]

\[
\bar{\kappa}_{xy} = -\bar{\kappa}_{yx} \approx \kappa_{xy} - \frac{4\pi^4 T^3 \omega_c \gamma^2}{d p_F^2 (\Gamma^2 + \omega_c^2)((\Gamma + \gamma)^2 + \omega_c^2)}.
\]

The thermoelectric conductivity is given by

\[
\alpha_{xx} = \alpha_{yy} = -\frac{e\nu \pi^2 T v_F (\Gamma + \gamma)(ap_F + v_F(\Gamma - \Gamma_1 p_F + 2\gamma)) + \omega_c^2 (3v_F + \Gamma_1 p_F v_F - a \Gamma p_F)}{p_F (\Gamma^2 + \omega_c^2)((\Gamma + \gamma)^2 + \omega_c^2)},
\]

18
\[
\alpha_{xy} = \alpha_{yx} = \frac{e\nu \pi^2 T}{d^3} \frac{p_F(a \Gamma - \Gamma_1 v_F)(2 \Gamma + \gamma) + v_F(2\gamma^2 + 2\omega_c^2 + 3\Gamma\gamma)}{p_F(\Gamma^2 + \omega_c^2)((\Gamma + \gamma)^2 + \omega_c^2)}.
\]

(87b)

We now unpack the thermal and thermoelectric conductivities. We first focus on the non-interacting limit \(\gamma = 0\). The Wiedemann-Franz law is obeyed component-wise:

\[
\bar{\kappa}_{ij} \approx \kappa_{ij} \approx \frac{\pi^2 T}{3e^2} \sigma_{ij},
\]

(88)
as is the Mott relation:

\[
\alpha_{ij} = -\frac{\pi^2 T}{3e} \frac{\partial \sigma_{ij}}{\partial \mu}.
\]

(89)

This relation can be checked using methods analogous to (57). In the limit of large \(\gamma\), we find that

\[
\left( \begin{array}{cc} \sigma_{xx} & T\alpha_{xx} \\ T\alpha_{xx} & T\kappa_{xx} \end{array} \right) = \frac{1}{\mathcal{M}} \left( \begin{array}{cc} e^2n^2 & -eTs \nu \left( T^2 \right) \\ -eTs \nu \left( T^2 \right) & (Ts)^2 + \omega_c^2 \end{array} \right) \frac{\Gamma}{\Gamma^2 + \omega_c^2} = \left( \begin{array}{cc} e^2n^2 & -eTs \nu \left( T^2 \right) \\ -eTs \nu \left( T^2 \right) & (Ts)^2 \end{array} \right) \frac{\mathcal{M}\Gamma}{(\mathcal{M}\Gamma)^2 + (enB)^2}.
\]

(90a)

\[
\left( \begin{array}{cc} \sigma_{xy} & T\alpha_{xy} \\ T\alpha_{xy} & T\kappa_{xy} \end{array} \right) = \frac{1}{\mathcal{M}} \left( \begin{array}{cc} e^2n^2 & -eTs \nu \left( T^2 \right) \\ -eTs \nu \left( T^2 \right) & -\omega_c \nu \left( T^2 \right) + \omega_c^2 \end{array} \right) \frac{-\omega_c}{\Gamma^2 + \omega_c^2} = \left( \begin{array}{cc} e^2n^2 & -eTs \nu \left( T^2 \right) \\ -eTs \nu \left( T^2 \right) & (Ts)^2 \end{array} \right) \frac{-enB}{(\mathcal{M}\Gamma)^2 + (enB)^2}.
\]

(90b)

These latter equations are consistent with the predictions of the memory matrix formalism at weak disorder and weak magnetic field strength [37].

When \(\Gamma, \gamma\), and \(\omega_c\) are all comparable, we conclude from (85) and (87) that the Wiedemann-Franz and Mott relations are no longer valid. The thermal conductivity depends rather simply on interactions: to leading order in \(T/T_F\), we may simply replace \(\Gamma \rightarrow \Gamma + \gamma\), as in Section 4.3. However, the thermoelectric conductivity \(\alpha_{ij}\) is far more complicated, and we do not see a simple way to disentangle the effects of finite magnetic field, disorder and interaction strength.

6 Discussion and Conclusions

The WF and Mott laws are strongly affected by electron-electron interactions in the hydrodynamic regime entirely within the Fermi liquid paradigm. Even an arbitrary suppression of the WF ratio well below the ideal Lorenz number does not necessarily signify any NFL, but may only indicate strong interaction effects in the hydrodynamic regime [11]. The fact that most normal 3D metals manifest the ideal WF behavior is a consequence of the fact that \(\Gamma \gg \gamma\) by virtue of the large \(T_F \sim 10^4-10^5\) K, and also the relative strength of electron-phonon scattering. This is in spite of 3D normal metals being ‘strongly interacting’ in the sense of having a large dimensionless coupling constant \(r_s \sim 6\). At low temperatures, where electron-phonon interaction is negligible compared with electron-electron interaction, the electron-impurity scattering is stronger than electron-electron interaction. We do not see any obvious theoretical obstruction to a 3D metal where electronic hydrodynamics occurs so long as the condition \(\gamma > \Gamma\) exists; a recent experiment suggests such 3D systems exist [20].

We expect that at least some of the experimentally observed strange metals are in a hydrodynamic regime. Earlier discussion of this point may also be found in [13, 14]. Is it also possible that this hydrodynamic regime is reasonably described by the Fermi liquid approach detailed in this paper? Answering this question requires careful material-specific considerations of each system to investigate (i) whether \(\gamma \gg \Gamma\) is satisfied, and (ii) if the failure of the WF law is happening in a temperature range consistent with our Eq. (6). Such an investigation is well beyond the scope of the current work, but is worth future consideration. Perhaps (at least in some portions of an often complicated phase diagram) strange metals
are 'strange' only in the sense of being hydrodynamic metals. Indeed, most strange metals have rather low $T_F \sim 10^3$ K, so strong interaction effects could easily drive the system into the hydrodynamic regime. In many strange metals, the large $T$-linear resistivity, violating the Mott-Ioffe-Regel 'bound' [59], is used as additional evidence for NFL physics [1, 4]. The validity of our kinetic approach is suspect in this limit. Nevertheless, building upon [14], we propose a more careful study of thermal and electrical transport in the Fermi liquid regime of strange metals, where “conventional” $T^2$ resistivity is observed just outside of the “critical fan” of $T$-linear resistivity, and where the kinetic theory of transport should apply. It is reasonable to assume that disorder has the same origin in both strange and conventional portions of the phase diagram [14], so we expect that a clear understanding of the roles of umklapp, phonon scattering, and hydrodynamic effects in this regime will shed light on the origin of $T$-linear resistivity.

All solid state materials have (at least) three distinct scattering mechanisms affecting transport: electron-impurity scattering (controlling low-temperature transport), electron-phonon scattering (controlling high-temperature transport), and electron-electron scattering (controlling the ballistic to hydrodynamic crossover, in the absence of umklapp). Hydrodynamics can only be observed if $\gamma \sim T^2/T_F$ is the dominant of these three scattering mechanisms. This rules out hydrodynamic observations at high temperatures, since electron-phonon scattering rates (not considered in our work) scaling as $\sim T/T_D$ (for $T > T_D$), where $T_D (\sim 10^2 - 10^3$ K) is the typical phonon energy scale in most materials, is the dominant resistive scattering mechanism in all electronic materials at higher temperatures. At low temperatures, the requirement $\gamma >> \Gamma$ necessitates both a very pure system so that the impurity disorder strength is weak and also a Fermi temperature low enough for $\gamma$ not to be too small. For a Coulomb-interacting Fermi liquid with parabolic band dispersion with an effective mass $m$, the requirement for hydrodynamics becomes $m/n^{2/d}$ being very large, where $n$ is the carrier density and $d$ the spatial dimension. In addition, the "low-temperature" condition $T \ll T_D$ must be satisfied in order for phonon scattering to be unimportant. It turns out, as emphasized already, this quantity is rather small in 3D metals by virtue of $n$ being too large. However, in 2D GaAs systems, the hydrodynamic condition can easily be satisfied, both because the carrier density can be made very low and the impurity disorder can be made very small by utilizing modulation doping.

Indeed, the systems where we believe that our hydrodynamic predictions are most likely going to be verified are low-disorder, high-mobility 2D systems such as modulation-doped 2D GaAs systems and high-quality graphene layers. Converting the available experimental 2D mobility values and theoretical (but experimentally verified) electron-electron scattering strengths to effective scattering times, we find that the condition $\gamma >> \Gamma$ is obeyed in 2D GaAs structures (both n-doped and p-doped) down to 1 K or below in dilute systems of density $\sim 10^{11}$ cm$^{-2}$ [60, 61]. At these low temperatures, phonon effects are completely negligible, and therefore, our approximations should apply uncritically with the dilute 2D GaAs system being deep in the hydrodynamic regime. We predict a strong failure of the WF law in these systems, and our prediction of (6) can be directly verified by varying the temperature. To be specific, in the 2D n-GaAs system (assuming a mobility of $10^7$ cm$^2$/V·s), the electron-electron and the electron-impurity scattering times at $T = 3$ K and $n = 3 \times 10^{11}$ cm$^{-2}$ are respectively $\tau_{ee} = 1/\gamma = 20$ ps and $\tau_{ei} = 1/\Gamma = 400$ ps, making $\gamma \sim 20\Gamma$, thus placing the system deep into the hydrodynamic regime. Lowering $T$ to 100 mK in the same sample hardly modifies the mobility (i.e. $\Gamma$), but $\tau_{ee}$ increases to 25000 ps, driving the system to $\gamma \ll \Gamma$. Since $T_F = 125$ K for $n = 3 \times 10^{11}$ cm$^{-2}$, the low-temperature condition necessary for the applicability of our Eq. (6) remains valid throughout. Thus, changing temperature from 3 K to 100 mK in a high mobility 2D n-GaAs system would be an ideal experiment for the verification of our theory of the failure of the WF law. The same is even more true for 2D p-GaAs holes also, since the hole mass is larger than the electron mass in GaAs, thus making $\gamma$ effectively larger at the same temperature since

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6We cannot formally rule out momentum-conserving electron-phonon scattering as playing an important role, but this seems unlikely in most materials.
In both cases, electron-phonon interaction is at least an order of magnitude weaker than even the electron-impurity interaction at these cryogenic temperatures, making the high-mobility 2D GaAs structures the ideal systems for studying hydrodynamic transport effects.

Unfortunately, we find that the other well-known 2D semiconductor system, namely Si MOSFETs [62], does not satisfy the hydrodynamic condition (it comes close, but falls below the minimal requirement of $\gamma \gtrsim \Gamma$) in any accessible density or temperature range. Even the purest 2D Si system is simply not pure enough, at present.

In graphene, one has to go rather close to the Dirac point ($n \sim 10^9 \text{ cm}^{-2}$) to achieve the hydrodynamic constraint at $T \sim 50$ K. Here phonon effects are weak, but not negligible. One therefore requires extremely clean samples so that impurity-induced puddle effects do not overwhelm the hydrodynamic effects at such low densities. Such experiments have been performed and interpreted based on hydrodynamic theories recently [27, 31], but more work should be done to directly verify our predictions. Recently, the relevance of clean bilayer graphene in the context of 2D hydrodynamic transport has been pointed out [63].

In this paper, we have focused on the limit $T \ll T_F$. Assuming reasonable forms for the electron-electron interactions, in this limit we rigorously reduced the calculation of thermoelectric conductivities to a finite-dimensional linear algebra problem. We emphasize, however, that the techniques described in this paper immediately and straightforwardly generalize to $T \sim T_F$ (so long as long-lived quasiparticles exist), and to systems without rotational invariance. In these more complicated settings, it is likely that the evaluation of the thermoelectric conductivity matrix must be done numerically, following the theoretical formalism described in this work.

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### Low Temperature Expansion

In this appendix, we describe more explicit details about the low temperature expansion of inner products in the $|q^n\rangle$ and $|n\rangle$ bases, in Section 4. We employ the integral

$$
\int_{-\infty}^{\infty} \frac{dx}{(1 + e^x)^n} e^{(m+1)x} = \frac{\Gamma(1 + m)\Gamma(n - m - 1)}{\Gamma(n)} = F(m, n),
$$

(91)

Ignoring the singularity at the origin of momentum space only leads to exponentially small corrections ($e^{-T_F/T}$) in our theory. We then define

$$
G(m, n, p) \equiv \frac{\partial^p F(m, n)}{\partial m^p} = \int_{-\infty}^{\infty} \frac{dx}{(1 + e^x)^n} e^{(m+1)x} x^p.
$$

(92)

Having defined $G(m, n, p)$, all integrals over momentum space in the inner products can be analytically evaluated. But, the process is a little bit subtle. In particular, observe that the derivative of the Fermi function must be carefully expanded:

$$
\frac{e^{(q)/T}}{T(1 + e^{(q)/T})^2} = \frac{e^{vFq/T}}{(1 + e^{vFq/T})^2} \left[ 1 - \frac{(e^{vFq/T} - 1)}{(1 + e^{vFq/T})^2} \left( \frac{ap^2}{2} + \frac{bp^3}{3} \right) + -\frac{(e^{2vFq/T} - 4e^{vFq/T} + 1)}{2(1 + e^{vFq/T})^2} \left( \frac{ap^2}{2} + \frac{bp^3}{3} \right)^2 + \cdots \right]
$$

(93)
This means that
\[
\langle q^n | q^m \rangle \approx \frac{\nu}{d} \int_{-\infty}^{\infty} dq \left( 1 + \frac{q}{p_F} \right)^{d-1} \frac{e^{\epsilon(q)/T}}{T(1 + e^{\epsilon(q)/T})^2} q^{m+n}.
\] (94)
is not straightforwardly given by something proportional to \( G(0,2,m+n) \). In \( d = 2 \), for example, we find the complicated Taylor series
\[
\langle q^n | q^m \rangle \approx \frac{\nu}{d} \int_{-\infty}^{\infty} dq \left[ G(0,2,m+n) - \frac{aT}{2v_F^2} (G(1,3,m+n+2) - G(0,3,m+n+2)) \\
- \frac{bT^2}{3v_F^2} (G(1,3,m+n+3) - G(0,3,m+n+3)) \\
+ \frac{a^2T^2}{8v_F^2} (G(2,4,m+n+4) - 4G(1,4,m+n+4) + G(0,4,m+n+4)) \\
+ \frac{T}{v_F p_F} \left( G(0,2,m+n+1) - \frac{aT}{2v_F^2} (G(1,3,m+n+3) - G(0,3,m+n+3)) \right) \right] + O(T^3). \] (95)
The expansion of the inner product to \( O(T^2) \), as above, will be adequate for the purposes of this paper.

### B Thermodynamic Identities

In this appendix, we describe the relationship between entropy, charge and the inner products in our kinetic theory.

First we re-derive an identity observed in [38]:
\[
\langle J_x | P_x \rangle = \int \frac{d^d p}{(2\pi \hbar)^d} \left( -\frac{\partial f_{eq}}{\partial \epsilon} \right) v_x p_x = \int \frac{d^d p}{(2\pi \hbar)^d} \left( -\frac{\partial f_{eq}}{\partial p_x} \right) p_x = \int \frac{d^d p}{(2\pi \hbar)^d} f_{eq} = n. \] (96)

Observe that the zero temperature charge density is given by
\[
n_0 = \frac{p_F^d}{d(2\pi \hbar)^d} \Omega_d, \] (97)
where \( \Omega_d \) denotes the volume of the unit sphere in \( d \) spatial dimensions. Secondly, the density of states is given by
\[
\nu = \frac{\partial n_0}{\partial \mu} = \frac{1}{v_F} \frac{\partial n_0}{\partial p_F} = \Omega_d \frac{p_F^d - 1}{v_F (2\pi \hbar)^d}. \] (98)

In the large Fermi surface limit, in \( d = 2 \), we have
\[
n \approx \frac{\nu v_F p_F^2}{2}. \] (99)

Next, we find
\[
\langle Q_x | P_x \rangle = \int \frac{d^d p}{(2\pi \hbar)^d} \left( -\frac{\partial f_{eq}}{\partial \epsilon} \right) v_x \epsilon p_x = \int \frac{d^d p}{(2\pi \hbar)^d} \left( -\frac{\partial f_{eq}}{\partial p_x} \right) p_x \epsilon = \int \frac{d^d p}{(2\pi \hbar)^d} f_{eq} (\epsilon + p_x v_x). \] (100)
The pressure is given by
\[
P = \int \frac{d^d p}{(2\pi \hbar)^d} T \log \left( 1 + e^{-(\epsilon - \mu)/T} \right) \] (101)
and we find (using the same integration by parts tricks as before)

\[
\int \frac{d^d p}{(2\pi \hbar)^d} f_{eq}p_x v_x = \int \frac{d^d p}{(2\pi \hbar)^d} T \log \left(1 + e^{-\epsilon/T}\right).
\]  

(102)

We conclude that

\[
\langle Q_x | P_x \rangle = \langle \epsilon \rangle + P = Ts,
\]

(103)

where \( s \) is the entropy density. In the last step, we have used the Gibbs-Duhem identity together with the fact that we have fixed \( \mu = 0 \) by our conventions. In a Fermi liquid, we also have

\[
s = \left(\frac{\partial P}{\partial T}\right)_\mu \approx \frac{\pi^2 T^3}{3} \nu.
\]

(104)

Finally, the momentum-momentum susceptibility is given by

\[
\mathcal{M} = \lim_{v_x \to 0} \frac{1}{v_x} \int \frac{d^d p}{(2\pi \hbar)^d} p_x f_{eq}(\epsilon - p_x v_x) = \int \frac{d^d p}{(2\pi \hbar)^d} \frac{p^2}{d} \left(-\frac{\partial f_{eq}}{\partial \epsilon}\right) \approx \frac{\nu p_F^2}{d}.
\]

(105)

Note that (in \( d = 2 \), at least)

\[
n \frac{\mathcal{M}}{v_F} = \frac{p_F}{p_F}.
\]

(106)

This identity allows one to show (90).

\section{The First Three Basis Vectors}

We must keep track of \( |0\rangle \), \( |1\rangle \) and \( |2\rangle \), as well as keep track of \( O(T^2) \) corrections to \( W_{\text{imp}} \), in order to correctly compute (68). The vectors

\[
|J_x\rangle \approx \left(v_F - \frac{\pi^2 T^2}{6p_F^2 v_F^3}\right) |0\rangle + \frac{\pi T}{\sqrt{3} p_F} |1\rangle,
\]

(107a)

\[
|Q_x\rangle \approx \left(\frac{\pi^2 T^2}{3p_F} + \frac{21\pi^4 T^4}{8p_F^2 v_F^2}\right) |0\rangle + \left(\frac{\pi T v_F}{\sqrt{3}} - \frac{31\pi^3 T^3}{15\sqrt{3} p_F^2 v_F^2}\right) |1\rangle + \left(\frac{2\pi^2 T^2}{\sqrt{5} p_F^2} + \frac{14\pi^4 T^4}{15\sqrt{5} p_F^2 v_F^2}\right) |2\rangle.
\]

(107b)

We have included the first non-trivial subleading corrections in \( T \), which will be necessary to compute the first non-trivial subleading corrections to conductivity. As noted in Section 4.4, we have assumed a quadratic dispersion relation; hence \( 2T_F = p_F v_F \), and \( a p_F = v_F \). Using (40) and the fact from (32) that \( A_{\text{imp}} \) is a constant, we obtain

\[
\frac{\Gamma_1}{\Gamma} = -\frac{2}{p_F},
\]

(108a)

\[
\frac{\Gamma_2}{\Gamma} = \frac{3}{p_F}.
\]

(108b)

Finally,

\[
W_{\text{imp}} \approx \Gamma \left(1 + \frac{5\pi^2 T^2}{8T_F^2}\right) |0\rangle \langle 0| - \frac{\sqrt{3} \pi T}{2T_F} (|1\rangle \langle 0| + |0\rangle \langle 1|) + \Gamma \left(1 + \frac{137\pi^2 T^2}{40T_F^2}\right) |1\rangle \langle 1| + \frac{2\pi^2 T^2}{\sqrt{5} T_F^2} (|2\rangle \langle 0| + |0\rangle \langle 2|)
\]

\[
- \sqrt{\frac{12}{5}} \frac{\pi T}{T_F} (|2\rangle \langle 1| + |1\rangle \langle 2|) + \Gamma \left(1 + \frac{2523\pi^2 T^2}{280T_F^2}\right) |2\rangle \langle 2|.
\]

(109)
Employing (23), we find (68).

Finally, let us justify that the relaxation time approximation for electron-electron collisions does not change the form of (68). Using (107b), the electron-electron contribution to \( T \bar{\kappa} \) is

\[
T \bar{\kappa} \approx \frac{\pi^2 T^2 v_F^2}{3} \left( 1 - \frac{1}{\Gamma} \langle 1 | (\bar{W} - \Gamma) | 1 \rangle + \cdots \right) = \frac{\pi^2 T^2 v_F^2}{3} \left[ 1 - \frac{\langle 1 | W | 1 \rangle}{\Gamma} + \cdots \right].
\]

As \( \langle 1 | W | 1 \rangle \) is \( O(T^2) \), we conclude that so long as \( \alpha^2 T^2 / T_F \equiv \langle 1 | W | 1 \rangle, (68) \) is unchanged for arbitrary collision integrals.

**D Thermal Conductivity in the Hydrodynamic Limit**

In this appendix we explicitly compute the hydrodynamic dissipative coefficient \( \kappa_q \). In a Galilean-invariant fluid, the heat current is given by

\[
Q = -T s v_{\text{hydro}} - \kappa_q \nabla T. \tag{111}
\]

To see how this arises within kinetic theory, consider the following component of the time-independent Boltzmann in a translation-invariant fluid:

\[
v_j \partial_j | Q_i \rangle \approx -\gamma \left( | Q_i \rangle - \frac{\langle Q_i | P_j \rangle}{\langle P | P \rangle} | P_j \rangle \right). \tag{112}
\]

In a hydrodynamic limit, \( v_j \partial_j | Q_i \rangle \) is dominated by any vector associated with a conserved quantity – since these gradients are long lived modes. Assume for simplicity that charge, energy and momentum are the only conserved quantities: the first two are given by

\[
| n \rangle = \int \frac{d^d p}{(2\pi \hbar)^d} | p \rangle,
\]

\[
| e \rangle = \int \frac{d^d p}{(2\pi \hbar)^d} \epsilon(p) | p \rangle. \tag{113b}
\]

Note the identities

\[
\langle n | n \rangle = \int \frac{d^d p}{(2\pi \hbar)^d} \left( -\frac{\partial f_{\text{eq}}}{\partial \epsilon} \right) = \frac{\partial n}{\partial \mu} \equiv \chi, \tag{114a}
\]

\[
\langle e | e \rangle = \int \frac{d^d p}{(2\pi \hbar)^d} \epsilon^2 \approx \frac{\pi^2 T^2 \nu}{3}, \tag{114b}
\]

\[
\langle n | e \rangle \sim T^2 \tag{114c}
\]

with \( \chi \) the charge compressibility; note \( \chi \sim T^0 \). Since

\[
\langle n | v_j | Q_i \rangle = \frac{\delta_{ij}}{d} \int \frac{d^d p}{(2\pi \hbar)^d} \left( -\frac{\partial f_{\text{eq}}}{\partial \epsilon} \right) v^2 \epsilon \sim T^2, \tag{115a}
\]

\[
\langle e | v_j | Q_i \rangle = \frac{\delta_{ij}}{d} \int \frac{d^d p}{(2\pi \hbar)^d} \left( -\frac{\partial f_{\text{eq}}}{\partial \epsilon} \right) v^2 \epsilon^2 \approx \langle e | e \rangle \frac{v_F^2}{d} \delta_{ij} \tag{115b}
\]

using (103), we conclude that

\[
| Q_i \rangle - T s \frac{| P_i \rangle}{\langle P | P \rangle} = -\frac{v_F^2}{d \gamma} \partial_i | e \rangle + O(T^2 \partial_i | n \rangle) \tag{116}
\]
In order to relate fluctuations in temperature $\delta T$ to $|\epsilon\rangle$, observe that

$$f_{eq}(T + \delta T) - f_{eq}(T) = \left( \frac{\epsilon}{T} \delta T \right) \left( -\frac{\partial f_{eq}}{\partial \epsilon} \right),$$

which implies that

$$\langle e|\Phi\rangle = \delta T \times \frac{\langle e|e\rangle}{T} = \delta T \times \frac{\pi^2 T \nu}{3}.$$  \hfill (118)

Taking the inner product of (116) with a generic distribution function $|\Phi\rangle$, and noting that the approximate low-temperature identity $J = n v_{\text{hydro}}$ implies that $\langle \Phi|P_i\rangle = \langle P|P\rangle v_{\text{hydro},i}$, we conclude that (111) holds with

$$\kappa_Q = \frac{\pi^2 \nu^2 T^2}{3d\gamma}.$$  \hfill (119)

This expression precisely agrees with the open-circuit thermal conductivity (67) in the hydrodynamic limit $\gamma \gg \Gamma$.

### References

[1] J. A. N. Bruin, H. Sakai, R. S. Perry, and A. P. Mackenzie. “Similarity of scattering rates in metals showing $T$-linear resistivity”, *Science* **339** 804 (2013).

[2] V. J. Emery and S. A. Kivelson. “Superconductivity in bad metals”, *Physical Review Letters* **74** 3253 (1995).

[3] R. Mahajan, M. Barkeshli, and S. A. Hartnoll. “Non-Fermi liquids and the Wiedemann-Franz law”, *Physical Review* **B88** 125107 (2013), arXiv:1304.4249.

[4] S. A. Hartnoll. “Theory of universal incoherent metallic transport”, *Nature Physics* **11** 54 (2015), arXiv:1405.3651.

[5] S. A. Hartnoll, A. Lucas, and S. Sachdev. “Holographic quantum matter”, arXiv:1612.07324.

[6] R. Franz and G. Wiedemann. “Ueber die Wärme-Leitungsfähigkeit der Metalle”, *Annalen der Physik* **165** 497 (1853).

[7] R. Lorenz. “Ueber das Leitungsvermögen der Metalle für Wärme und Electricität”, *Annalen der Physik* **249** 582 (1881).

[8] J. Ziman. *Electrons and Phonons* (Oxford University Press, 1960).

[9] G. S. Kumar, G. Prasad, and R. O. Pohl. “Experimental determinations of the Lorenz number”, *Journal of Materials Science* **28** 4261 (1993).

[10] R. W. Hill, C. Proust, L. Taillefer, P. Fournier, and R. L. Greene. “Breakdown of Fermi-liquid theory in a copper-oxide superconductor”, *Nature* **414** 711 (2001).

[11] A. Principi and G. Vignale. “Violation of the Wiedemann-Franz law in hydrodynamic electron liquids”, *Physical Review Letters* **115** 056603 (2015).

[12] S. Sachdev. *Quantum Phase Transitions* (Cambridge University Press, 2nd ed., 2011).
[13] R. A. Davison, K. Schalm, and J. Zaanen. “Holographic duality and the resistivity of strange metals”, Physical Review B89 245116 (2014), arXiv:1311.2451.

[14] A. Lucas and S. A. Hartnoll. “Resistivity bound for hydrodynamic bad metals”, Proceedings of the National Academy of Sciences 114 11344 (2017), arXiv:1704.07384.

[15] A. Lucas and K. C. Fong. “Hydrodynamics of electrons in graphene”, Journal of Physics: Condensed Matter 30 053001 (2018), arXiv:1710.08425.

[16] E. H. Hwang and S. Das Sarma. “Limit to two-dimensional mobility in modulation-doped GaAs quantum structures: how to achieve a mobility of 100 million”, Physical Review B77 235437 (2008), arXiv:0804.4684.

[17] D. A. Bandurin et al. “Negative local resistance due to viscous electron backflow in graphene”, Science 351 1055 (2016), arXiv:1509.04165.

[18] R. Krishna Kumar et al. “Super-ballistic flow of viscous electron fluid through graphene constric-
tions”, Nature Physics 13 1182 (2017), arXiv:1703.06672.

[19] P. J. W. Moll, P. Kushwaha, N. Nandi, B. Schmidt, and A. P. Mackenzie. “Evidence for hydrodynamic electron flow in PdCoO$_2$”, Science 351 1061 (2016), arXiv:1509.05691.

[20] J. Gooth, F. Menges, C. Shekhar, V. Süß, N. Kumar, Y. Sun, U. Drechsler, R. Zierold, C. Felser, and B. Gotsmann. “Electrical and thermal transport at the Planckian bound of dissipation in the hydrodynamic electron fluid of WP$_2$”, arXiv:1706.05925.

[21] E. V. Levinson, G. M. Gusev, A. D. Levin, E. V. Levinson, and A. K. Bakarov. “Viscous electron flow in mesoscopic two-dimensional electron gas”, AIP Advances 8 025318 (2018), arXiv:1802.09619.

[22] X. Lin, B. Fauque, and K. Behnia. “Scalable $T^2$ resistivity in a small single-component Fermi surface”, Science 349 945 (2015), arXiv:1508.07812.

[23] E. Mikheev, S. Raghavan, J. Y. Zhang, P. B. Marshall, A. P. Kajdos, L. Balents, and S. Stemmer. “Carrier density independent scattering rate in SrTiO$_3$-based electron liquids”, Scientific Reports 6 20865 (2016), arXiv:1512.02294.

[24] A. V. Andreev, S. A. Kivelson, and B. Spivak. “Hydrodynamic description of transport in strongly correlated electron systems”, Physical Review Letters 106 256804 (2011), arXiv:1011.3068.

[25] I. Torre, A. Tomadin, A. K. Geim, and M. Polini. “Non-local transport and the hydrodynamic shear viscosity in graphene”, Physical Review B92 165433 (2016), arXiv:1508.00363.

[26] L. Levitov and G. Falkovich. “Electron viscosity, current vortices and negative nonlocal resistance in graphene”, Nature Physics 12 672 (2016), arXiv:1508.00836.

[27] A. Lucas, J. Crossno, K. C. Fong, P. Kim, and S. Sachdev. “Transport in inhomogeneous quantum critical fluids and in the Dirac fluid in graphene”, Physical Review B93 075426 (2016), arXiv:1510.01738.

[28] H. Guo, E. Ilseven, G. Falkovich, and L. Levitov. “Higher-than-ballistic conduction of viscous electron flows”, Proceedings of the National Academy of Sciences 114 3068 (2017), arXiv:1607.07269.

[29] F. M. D. Pellegrino, I. Torre, A. K. Geim, and M. Polini. “Electron hydrodynamics dilemma: whirlpools or no whirlpools”, Physical Review B94 155414 (2016), arXiv:1607.03726.
T. Scaffidi, N. Nandi, B. Schmidt, A. P. Mackenzie, and J. E. Moore. “Hydrodynamic electron flow and Hall viscosity”, Physical Review Letters 118 226601 (2017), arXiv:1703.07325.

J. Crossno et al. “Observation of the Dirac fluid and the breakdown of the Wiedemann-Franz law in graphene”, Science 351 1058 (2016), arXiv:1509.04713.

F. Ghahari, H-Y. Xie, T. Taniguchi, K. Watanabe, M. S. Foster, and P. Kim. “Enhanced thermoelectric power in graphene: violation of the Mott relation by inelastic scattering”, Physical Review Letters 116 136802 (2016), arXiv:1601.05859.

R. P. Smith, M. Sutherland, G. G. Lonzarich, S. S. Saxena, N. Kimura, S. Takashima, M. Nohara, and H. Takagi. “Marginal breakdown of the Fermi-liquid state on the border of metallic ferromagnetism”, Nature 455 1220 (2008).

H. Pfau et al. “Thermal and electrical transport across a magnetic quantum critical point”, Nature 484 493 (2013), arXiv:1307.1066.

S. A. Hartnoll, P. K. Kovtun, M. Müller, and S. Sachdev. “Theory of the Nernst effect near quantum phase transitions in condensed matter, and in dyonic black holes”, Physical Review B76 144502 (2007), arXiv:0706.3215.

A. Lucas and S. A. Hartnoll. “Kinetic theory of transport for inhomogeneous electron fluids”, Physical Review B97 045105 (2018), arXiv:1706.04621.

A. Lucas. “Kinetic theory of electronic transport in random magnetic fields”, Physical Review Letters 120 116603 (2018), arXiv:1710.11141.

K. C. Fong and K. C. Schwab. “Ultrasensitive and wide-bandwidth thermal measurements of graphene at low temperatures”, Physical Review X2 031006 (2012), arXiv:1202.5737.

J. Crossno, X. Liu, T. A. Ohki, P. Kim, and K. C. Fong. “Development of high frequency and wide bandwidth Johnson noise thermometry”, Applied Physics Letters 106 023121 (2015), arXiv:1411.4596.

P. Ledwith, H. Guo, and L. Levitov. “Fermion collisions in two dimensions”, arXiv:1708.01915.

S. Das Sarma and E. H. Hwang. “Charged impurity scattering limited low temperature resistivity of low density silicon inversion layers”, Physical Review Letters 83 164 (1999), arXiv:cond-mat/9812216.

S. Das Sarma and E. H. Hwang. “Metallicity and its low-temperature behavior in dilute two-dimensional carrier systems”, Physical Review B69 195305 (2004), arXiv:cond-mat/0302047.

P. L. Bhatnagar, E. P. Gross, and M. Krook. “A model for collision processes in gases. I. Small amplitude processes in charged and neutral one-component systems”, Physical Review 94 511 (1954).

C. Hodges, H. Smith, and J. W. Wilkins. “Effect of Fermi surface geometry on electron-electron scattering”, Physical Review B4 302 (1971).
[47] G. F. Giuliani and J. J. Quinn. “Lifetime of a quasiparticle in a two-dimensional electron gas”, *Physical Review* B26 4421 (1982).

[48] T. Jungwirth and A. H. MacDonald. “Electron-electron interactions and two-dimensional-two-dimensional tunneling”, *Physical Review* B53 7403 (1996), arXiv:cond-mat/9603001.

[49] L. Zheng and S. Das Sarma. “Coulomb scattering lifetime of a two-dimensional electron gas”, *Physical Review* B53 9964 (1996), arXiv:cond-mat/9602066.

[50] B. N. Narozhny, G. Zala, and I. L. Aleiner. “Interaction corrections at intermediate temperatures: dephasing time”, *Physical Review* B65 180202 (2002), arXiv:cond-mat/0201379.

[51] Q. Li and S. Das Sarma. “Finite temperature inelastic mean free path and quasiparticle lifetime in graphene”, *Physical Review* B87 085406 (2013), arXiv:1211.6430.

[52] S. Q. Murphy, J. P. Eisenstein, L. N. Pfeiffer, and K. W. West. “Lifetime of two-dimensional electrons measured by tunneling spectroscopy”, *Physical Review* B52 14825 (1995).

[53] H. Noh, M. P. Lilly, D. C. Tsui, J. A. Simmons, E. H. Hwang, S. Das Sarma, L. N. Pfeiffer, and K. W. West. “Interaction corrections to two-dimensional hole transport in the large-\(r_s\) limit”, *Physical Review* B68 165308 (2003), arXiv:cond-mat/0206519.

[54] M. Müller, L. Fritz, and S. Sachdev. “Quantum-critical relativistic magnetotransport in graphene”, *Physical Review* B78 115406 (2008), arXiv:0805.1413.

[55] H-Y. Xie and M. S. Foster. “Transport coefficients of graphene: Interplay of impurity scattering, Coulomb interaction, and optical phonons”, *Physical Review* B93 195103 (2016), arXiv:1601.05862.

[56] S. Das Sarma and E. H. Hwang. “Screening and transport in 2D semiconductor systems at low temperatures”, *Scientific Reports* 5 16655 (2015).

[57] F. Stern. “Calculated temperature dependence of mobility in silicon inversion layers”, *Physical Review Letters* 44 1469 (1986).

[58] S. Das Sarma. “Theory of finite-temperature screening in a disordered two-dimensional electron gas”, *Physical Review* B33 5401 (1986).

[59] N. E. Hussey, K. Takenaka, and H. Takagi. “Universality of the Mott-Ioffe-Regel limit in metals”, *Philosophical Magazine* B4 2847 (2004), arXiv:cond-mat/0404263.

[60] M. P. Lilly, J. L. Reno, J. A. Simmons, I. B. Spielman, J. P. Eisenstein, L. N. Pfeiffer, K. W. West, E. H. Hwang, and S. Das Sarma. “Resistivity of dilute 2D electrons in an undoped GaAs heterostructure”, *Physical Review Letters* 90 056806 (2003).

[61] M. J. Manfra, E. H. Hwang, S. Das Sarma, L. N. Pfeiffer, K. W. West, and A. M. Sergent. “Transport and percolation in a low-density high-mobility two-dimensional hole system”, *Physical Review Letters* 99 236402 (2007), arXiv:0710.3542.

[62] B. Hu, M. M. Yazdanpanah, B. E. Kane, E. H. Hwang, and S. Das Sarma. “Strongly metallic electron and hole 2D transport in an ambipolar Si-vacuum field effect transistor”, *Physical Review Letters* 115 036801 (2015), arXiv:1502.02956.
[63] D. Y. H. Ho, I. Yudhistra, N. Chakraborty, and S. Adam. “Theoretical determination of hydrodynamic window in monolayer and bilayer graphene from scattering rates”, Physical Review B97 121404 (2018), arXiv:1710.10272.