Resistivity of the two-dimensional Bose-Hubbard model at weak coupling

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We calculate the weak-coupling resistivity of the two-dimensional Bose Hubbard model, comparing with the more familiar fermionic case. At high temperature the resistivity is linear in T , while in the low-temperature normal state it is exponentially suppressed. We explore the density dependence and calculate the momentum relaxation rate.

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I. INTRODUCTION

Transport is the most fundamental probe in solid-state physics. Recently, new techniques have been developed to study the transport properties of quantum degenerate atomic gases $[1-3]$, giving us the ability to use ultracold atoms to expand our understanding. For example, many strongly correlated materials display unexplained temperature dependence of their resistivity [\[4–7\]](#page-4-0), and cold-atom experiments may reveal the underlying mechanisms. Cold-atom experiments are also exploring the rich behavior of transport in inhomogeneous settings such as quantum point contacts [\[8\]](#page-4-0). Beyond these solid-state-inspired studies, we can use atomic gases to explore completely novel transport regimes. Here we calculate the resistivity of a two-dimensional (2D) Bose gas in an optical lattice. As is familiar from other systems with bounded spectra $[9-13]$, we find that the high-temperature resistivity scales linearly in temperature, while the low-temperature behavior is more complex. The same techniques which were used to measure the resistivity of a 2D Fermi gas could be applied here to experimentally verify these results.

In most materials, resistivity is dominated by either electron-phonon scattering or electron-impurity scattering. Ultracold atoms have no phonons or impurities, so the only mechanism for momentum relaxation is the equivalent of electron-electron scattering [\[14\]](#page-4-0). Momentum is conserved in these processes, modulo a reciprocal lattice vector. Resistivity requires momentum relaxation and is thus related to the rate of umklapp scattering [\[15\]](#page-4-0), where the momentum of the particles changes by such a reciprocal lattice vector. We calculate this rate within the Born approximation, which works well at weak coupling, extracting the resistivity.

Resistivity is a quantity which is almost exclusively discussed in the context of fermions. Bosons, such as light, typically move ballistically. Moreover, there is no obvious optical equivalent of an electric field, making it challenging to even envision an optical transport experiment. Other common bosonic systems, such as phonons or magnons suffer these same challenges. Cold atoms, however, provide the perfect platform to extend transport measurements to bosons. All of the techniques that have been developed to probe transport in fermionic gasses can be used on bosonic atoms. Furthermore, the same mechanism which leads to the ohmic behavior of fermions also applies to bosons. Due to the lack of a Fermi surface, however, much of the phenomenology differs. Additionally, at very low temperatures a Bose gas becomes superfluid, with a vanishing resistivity. Our calculations only apply to the normal state.

At infinite temperature a lattice gas will have infinite resistivity [\[9,10,16\]](#page-4-0). All states are equally likely, and a uniform force does not change the momentum distribution or lead to currents. The leading corrections scale with 1/*T* , giving a *T* linear high-temperature resistivity. This argument is equally applicable to bosons and fermions.

For fermions [\[16\]](#page-4-0) at low temperature, all properties are dominated by excitations around the Fermi surface. Counting arguments then give that the umklapp contribution to resistivity generically scales as T^2 . Bosons are different. The low-temperature (but noncondensed) Bose distribution is peaked at small momentum. There are no allowed umklapp processes for these small momentum particles, so the resistivity is exponentially small.

At very low temperatures a 2D interacting Bose gas will undergo a Berezinskii-Kosterlitz-Thouless (BKT) transition to a superfluid state $[17-21]$, in which our calculation does not apply. The transition occurs at a temperature of order $T_{BKT} \sim nt/\ln(t/nU)$, where *n* is the average number of particles per site, t is the hopping matrix element and U is the on-site interaction strength. Typical densities are of the order of $0.1 < n < 2$.

We organize our paper as follows. In Sec. II we describe the 2D Bose-Hubbard model and our variational approach to solve the Boltzmann equation. In Sec. [III](#page-1-0) we present our results for the resistivity and its temperature dependence. In Sec. [IV](#page-2-0) we summarize and present some conclusions.

II. THE MODEL

We consider spinless bosonic particles on a square lattice, whose dynamics are governed by the Bose-Hubbard

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Hamiltonian [\[22\]](#page-4-0)

$$
H = -t \sum_{\langle i,j \rangle} b_i^{\dagger} b_j + \frac{U}{2} \sum_i b_i^{\dagger} b_i^{\dagger} b_i b_i
$$

$$
= \sum_{\mathbf{k}} \epsilon_{\mathbf{k}} b_{\mathbf{k}}^{\dagger} b_{\mathbf{k}} + \frac{U}{2N_s} \sum_{\mathbf{k}\mathbf{k}'\mathbf{q}} b_{\mathbf{k}}^{\dagger} b_{\mathbf{k}'}^{\dagger} b_{\mathbf{k}'-\mathbf{q}} b_{\mathbf{k}+\mathbf{q}}, \qquad (1)
$$

where b^{\dagger} and *b* are the creation and annihilation operators, respectively, *t* is the hopping amplitude, and *U* accounts for the local interaction. The single-particle dispersion relation is

$$
\epsilon_{\mathbf{k}} = -2t[\cos(k_x a) + \cos(k_y a)],\tag{2}
$$

where *a* is the lattice spacing, and the momentum space sums are taken over $k = (k_x, k_y)$, with $k_x, k_y = 2\pi n/L$. Here N_s is the number of lattice sites and $V = N_s a^2 = L^2$ is the volume (area).

We use a quantum kinetic approach [\[23,24\]](#page-4-0), describing the system in terms of the distribution function $f_k = f_k(\mathbf{r}, t)$, which measures how many particles have momentum **k** at position **r** and time *t*. We use the same symbol for the hopping matrix element and time, but the meaning should be clear from the context.

The distribution function obeys a Boltzmann equation:

$$
\frac{\partial f_{k}}{\partial t} + \mathbf{v}_{k} \cdot \nabla_{\mathbf{r}} f_{k} + \frac{e\mathbf{E}}{\hbar} \cdot \nabla_{k} f_{k} = I_{k}.
$$
 (3)

Here $\mathbf{v}_{\mathbf{k}} = \frac{1}{\hbar} \nabla_{\mathbf{k}} \epsilon_{\mathbf{k}}$. The electric field **E** is small, and we have taken the particles to have charge *e*. In the cold-atom context, the atoms are neutral and this designation is formal, since the only physical quantity is the gradient of the potential energy, $\nabla V = -eE$. The collision integral expresses the change in momentum occupation due to scattering,

$$
I_{\mathbf{k}} = \sum \left[\Omega_{\text{in}} - \Omega_{\text{out}} \right] \mathcal{W}_{\mathbf{k}\mathbf{k}'}^{\mathbf{k}'' \mathbf{k}''},\tag{4}
$$

where the sum is over \mathbf{k}' , \mathbf{k}'' , and \mathbf{k}''' in the first Brillouin zone $(-\pi/a < k_x, k_y < \pi/a)$. Quantum statistics are encoded in the factors $\Omega_{\text{in}} = f_{\mathbf{k}'''} f_{\mathbf{k}''}(1 + f_{\mathbf{k}'})(1 + f_{\mathbf{k}})$ and $\Omega_{\text{out}} = f_{\mathbf{k}} f_{\mathbf{k}} (1 + f_{\mathbf{k}''}) (1 + f_{\mathbf{k}'''})$. Using Fermi's golden rule, the transition rate $\mathcal{W}_{kk'}^{k''k'''} = \frac{2\pi}{\hbar} |\langle k''k'''| \mathcal{H} |kk' \rangle|^2 \delta(\epsilon_k + \epsilon_{k'} \epsilon_{\mathbf{k}''} - \epsilon_{\mathbf{k}'''}$) can be evaluated as

$$
\mathcal{W}_{\mathbf{k}\mathbf{k}'}^{\mathbf{k}''\mathbf{k}'''} = \frac{2\pi}{\hbar} \frac{U^2}{4N_s^2} \bar{\delta}_{\mathbf{k}+\mathbf{k}',\mathbf{k}''+\mathbf{k}'''} \delta(\epsilon_{\mathbf{k}} + \epsilon_{\mathbf{k}'} - \epsilon_{\mathbf{k}''} - \epsilon_{\mathbf{k}''}), \quad (5)
$$

where $\mathcal{H} = \frac{U}{2} \sum_i b_i^{\dagger} b_i^{\dagger} b_i b_i$ is the interaction Hamiltonian. Here $\bar{\delta}_{pq}$ is the periodic version of the Kronecker delta function, which is zero unless $\mathbf{p} - \mathbf{q}$ is a reciprocal lattice vector.

We linearize the Boltzmann equation by considering a small perturbation, $\delta f_{\mathbf{k}} = -\Phi_{\mathbf{k}}(\partial f_{\mathbf{k}}^0/\partial \epsilon_{\mathbf{k}})$, from the local equilibrium distribution function, $\hat{f}_{\mathbf{k}}^{0} = (e^{\beta(\epsilon_{\mathbf{k}}-\mu)} - 1)^{-1}$, writing

$$
f_{\mathbf{k}} = f_{\mathbf{k}}^0 - \Phi_k \frac{\partial f_{\mathbf{k}}^0}{\partial \epsilon_{\mathbf{k}}},
$$
 (6)

where the temperature $k_B T = \beta^{-1}$ and the chemical potential μ are uniform and time independent, implying $\nabla_{\mathbf{r}} f_{\mathbf{k}}^0 \approx 0$. The function $\Phi_{\mathbf{k}}$ is small and can be interpreted as a generalized current, generating a deviation from the equilibrium distribution. In a steady-state configuration $\partial_t f_{\mathbf{k}} = 0$, and the linearized Boltzmann equation becomes $e(\partial f_{\mathbf{k}}^0 / \partial \epsilon_{\mathbf{k}}) \mathbf{v}_{\mathbf{k}} \cdot \mathbf{E} =$ *I***k**, with

$$
I_{k} = -\beta \sum [\Phi_{k} + \Phi_{k'} - \Phi_{k''} - \Phi_{k'''}] \mathcal{P}_{kk'}^{k''k''}, \qquad (7)
$$

$$
\mathcal{P}_{\mathbf{k}\mathbf{k}'}^{\mathbf{k}''\mathbf{k}'''} = f_{\mathbf{k}}^0 f_{\mathbf{k}'}^0 (1 + f_{\mathbf{k}''}^0) (1 + f_{\mathbf{k}''}^0) \mathcal{W}_{\mathbf{k}\mathbf{k}'}^{\mathbf{k}''\mathbf{k}'''}. \tag{8}
$$

Following the approach in Refs. [\[25,26\]](#page-4-0), we make a variational ansatz, $\Phi_{\mathbf{k}} = \xi \phi_{\mathbf{k}}$, where $\phi_{\mathbf{k}} = (\nabla_{\mathbf{k}} \epsilon_{\mathbf{k}})_x = 2t \sin(k_x a)$ is fixed and ξ is a variational parameter. A more sophisticated ansatz would involve a linear combination of a set of linearly independent trial functions. If such a set is complete, then the solution is exact. We multiply the Boltzmann equation by −**k**/*V* and sum over **k**, resulting in

$$
\xi \mathbf{j} \cdot \mathbf{E} = \xi^2 P,\tag{9}
$$

where

$$
\mathbf{j} = -\frac{e}{V} \sum_{\mathbf{k}} \phi_{\mathbf{k}} \frac{\partial f_k^0}{\partial \epsilon_k} \mathbf{v}_{\mathbf{k}},\tag{10}
$$

$$
P = \frac{\beta}{4V} \sum [\phi_{\mathbf{k}} + \phi_{\mathbf{k}'} - \phi_{\mathbf{k}''} - \phi_{\mathbf{k}'''}]^2 \mathcal{P}_{\mathbf{k}\mathbf{k}'}^{\mathbf{k}''\mathbf{k}''}. \tag{11}
$$

Physically, these encode the charge density $J = \xi j$, and the entropy production due to collisions $\dot{S}_{\text{coll}} = \xi^2 P/T$. The connection between *P* and entropy comes from noting that the left-hand side of Eq. (9) is the power from ohmic dissipation $\mathbf{J} \cdot \mathbf{E} = -T \dot{S}_{\text{field}}$, which is converted into heat by collisions.

As argued in Appendix [C,](#page-4-0) $\xi = \mathbf{j} \cdot \mathbf{E}/P$ which satisfied Eq. (9) maximizes the entropy production and provides our best variational estimate. The resistivity is then

$$
\rho = \frac{P}{j^2},\tag{12}
$$

where $j = j_x$ is the *x* component of the current, parallel to the applied electric field. We also define a relaxation time τ from the Boltzmann equation according to $-\frac{\delta f_k}{\tau} = I_k$. Multiplying both sides by Φ_k/V and summing over **k**, the momentum relaxation rate $\Gamma = \tau^{-1}$ is

$$
\Gamma = \frac{ea}{\hbar} \frac{P}{j}.
$$
\n(13)

For a given temperature and density, *P* is an eightdimensional integral, but conservation of energy and momentum reduce it to five dimensions. The current j is a two-dimensional integral. In Appendix [A](#page-3-0) we explain how to efficiently calculate *P*. Details of the presented calculations are also addressed in the Supplemental Material [\[27\]](#page-4-0).

III. RESULTS

Our results are summarized by Fig. [1.](#page-2-0) Within the Born approximation, the resistivity scales with U^2 and has no other interaction dependence. The characteristic scale is $\rho_0 =$ $\frac{\hbar}{e^2}(\frac{U}{t})^2$. At high temperatures the resistivity is linear in *T*, approaching $\rho_{\infty} = 0.076(\frac{T}{t})\rho_0$. Such linear behavior is expected for any system with a bounded spectrum [\[9–13\]](#page-4-0).

The resistivity monotonically decreases with increasing density. This feature is related to the fact that the only momentum-changing collisions involve umklapp processes, $\mathbf{k}, \mathbf{k}' \rightarrow \mathbf{k}''$, \mathbf{k}''' , with $\mathbf{k} + \mathbf{k}' = \mathbf{k}'' + \mathbf{k}''' + \mathbf{Q}$, where **Q** is a

FIG. 1. Resistivity ρ as a function of temperature *T* for bosons (blue) or fermions (red) hopping on a 2D square lattice in the weakcoupling limit, $t \gg U$. Here $\rho_0 = \frac{\hbar}{e^2} (\frac{U}{t})^2$ is quadratic in the on-site interaction strength U . The hopping matrix element is t . The black line shows the classical limit, corresponding to vanishing density. The inset shows the bosonic data, on an inverse semilog scale. The thick yellow line has a slope of -4 , corresponding to $\rho \propto e^{-4t/T}$. At sufficiently low temperature, $T_{BKT} \sim nt/\ln(t/nU)$, the system will undergo a BKT transition to a superfluid, where this calculation is invalid. The shaded region indicates typical parameter values for the transition.

reciprocal lattice vector. If the energies $\epsilon_{\mathbf{k}}$ and $\epsilon_{\mathbf{k}'}$ are sufficiently small, then there are no energy-conserving processes of this form. Thus, low-energy particles have a smaller contribution to the resistivity than particles with high energy. For bosons, increasing the density will increase the fraction of particles with low energy, resulting in a suppressed resistivity.

Conversely, for fermions, increasing the density shifts the particle distribution to higher energy. Thus, the resistivity grows with density (below half-filling). In Fig. 1 we also show the fermion result, as calculated in Ref. [\[16\]](#page-4-0). At vanishing density the bosonic and fermionic results coincide, corresponding to a Boltzmann gas.

At low-temperature the resistivity is strongly suppressed, as exponentially small numbers of particles have sufficient energy to undergo umklapp processes. Thus, $\rho \sim e^{-\Delta/T}$. The inset in Fig. 1 illustrates this behavior by plotting $\ln(\rho/\rho_0)$ as a function of temperature 1/*T* . For all densities the data are well fit to $\Delta = 4t$, corresponding to the fact that in this regime the most important collisions occur between particles near the bottom of the band (with $\epsilon = -4t$) and particles at the van Hove singularity (with $\epsilon = 0$).

In addition to looking at the resistivity, it is useful to also explore the temperature dependence of the momentum relaxation rate Γ . While direct measures of Γ are often challenging in solid-state physics, cold-gas experiments have demonstrated the ability to measure it [\[28\]](#page-4-0). Figure 2 shows Γ versus T for bosons at different densities. The relaxation rate is a monotonically increasing function of density and temperature. More particles provide more opportunities for scattering, as well as contributing via Bose enhancement.

FIG. 2. Momentum relaxation rate Γ versus temperature T for bosons, at several different densities n . At high temperatures, Γ saturates at a value of $\Gamma_{\infty} = 0.152n(1 + n)U^2/\hbar t$. For low temperatures, the relaxation rate is exponentially suppressed.

Higher temperatures result in the occupation of larger momenta and, hence, more opportunities for umklapp scattering. At high temperatures the distribution function becomes flat and Γ saturates at $\Gamma_{\infty} = 0.152n(1+n)\Gamma_0$, with $\Gamma_0 = U^2/t\hbar$.

For low temperatures, Γ becomes exponentially suppressed, due to the low occupation of high-momentum modes. One also notes that the relaxation rate has only weak *n* dependence at low *T* . This is due to the fact that at sufficiently low temperatures, $\mu \rightarrow -4t$ for all densities. Thus, the occupation of high-momentum modes becomes density independent.

IV. SUMMARY AND CONCLUSIONS

Cold-atom experiments give us access to novel transport quantities, such as the resistivity of a Bose gas. Thus, we can elucidate the role of quantum statistics in transport.

Here we calculated the weak-coupling resistivity of a Bose gas in a 2D optical lattice. We find $\rho \propto T$ at high temperatures, similar to what was seen in experiments on fermionic atoms [\[1\]](#page-4-0). This illustrates that high-*T* linear resistivity is a property of the bound spectrum and is not solely a signature of exotic strongly correlated systems. In this high-temperature regime quantum statistics manifest in the density dependence of the resistivity. For bosons the resistivity falls with increasing density, while the opposite occurs for fermions.

It is somewhat counterintuitive that the relaxation rate saturates at high temperature, despite the diverging resistivity. This behavior can be traced to the properties of the effective mass. In a Drude picture, $\rho = m^*/ne^2 \tau$. For quantum particles in a lattice the mass is interpreted as the curvature of the dispersion, $(m^*)^{-1} = \partial_k^2 \epsilon_k$: Particles near the bottom of the band have a positive effective mass, while those at the top have a negative effective mass. At high temperature the atoms are evenly distributed throughout the Brillouin zone, and the appropriate average leads to *m*[∗] ∼ *T* [\[16\]](#page-4-0). This result can also be understood through the Nernst-Einstein relationship $\sigma = D\chi$, where *D* is the diffusion constant and χ is the compressibility.

The diffusion constant is related to the relaxation time and saturates at high temperature. The vanishing conductivity is attributed to a vanishing compressibility—as would be expected for any system with a bounded spectrum.

At low temperature we instead find that ρ is exponentially suppressed, as opposed to the fermionic $\rho \propto T^2$. This difference is due to the momentum distribution of the particles. At low temperatures the bosons predominantly occupy low *k* modes, which are incapable of umklapp scattering. Conversely, fermions form a Fermi sea. We also demonstrate that in the limit of vanishing density quantum statistics become irrelevant, and the fermionic and bosonic results coincide.

Experiments on bosons have largely focused on the superfluid regime. As this study illustrates, however, there are also many interesting phenomena which can be explored in the normal state. It would be very exciting to experimentally observe ohmic flow in a Bose gas. One could also explore the interplay between hydrodynamic and dissipative transport, as has been studied in recent solid-state experiments [\[29\]](#page-4-0).

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APPENDIX A: NUMERICALLY EVALUATING *P*

In the thermodynamic limit the discrete sums over momentum in Eq. [\(11\)](#page-1-0) become integrals and the periodic Kroenecker delta transforms into a sum of Dirac delta functions according to

$$
\frac{1}{V} \sum_{\mathbf{k}} \rightarrow \int \frac{d^2 \mathbf{k}}{(2\pi)^2}
$$

$$
V \overline{\delta}_{\mathbf{k} + \mathbf{k}', \mathbf{k}'' + \mathbf{k}'''} \rightarrow \sum_{Q} (2\pi)^2 \delta(\mathbf{k} + \mathbf{k}' - \mathbf{k}''' - \mathbf{k}''' - \mathbf{Q}), \quad (A1)
$$

where the reciprocal lattice vectors Q have components Q_x and Q_y which are integer multiples of $2\pi/a$. The collision term *P* is then

$$
P = \frac{U^2 a^4 \beta}{16\hbar (2\pi)^5} \sum_{Q} \int d^2 \mathbf{k} \int d^2 \mathbf{k}' \int d^2 \mathbf{k}'' \int d^2 \mathbf{k}'''
$$

× $(\phi_{\mathbf{k}} + \phi_{\mathbf{k}'} - \phi_{\mathbf{k}''} - \phi_{\mathbf{k}'''})^2 f_{\mathbf{k}}^0 f_{\mathbf{k}'}^0 (1 + f_{\mathbf{k}''}^0) (1 + f_{\mathbf{k}'''}^0)$
× $\delta(\mathbf{k} + \mathbf{k}' - \mathbf{k}'' - \mathbf{k}''' - \mathbf{Q}) \delta(\epsilon_{\mathbf{k}} + \epsilon_{\mathbf{k}'} - \epsilon_{\mathbf{k}''} - \epsilon_{\mathbf{k}'''}),$
(A2)

where, as already introduced, $\phi_{\mathbf{k}} = 2t \sin(k_x a)$, $\epsilon_{\mathbf{k}} =$ $-2t[\cos(k_x a) + \cos(k_y a)]$, and $f_k = (e^{\beta(\epsilon_k - \mu)} - 1)^{-1}$. Even after using the delta functions to eliminate integration variables, this is a five-dimensional integral. To more efficiently evaluate it, we change to center-of-mass and relative coordinates, writing

$$
\delta(\mathbf{k} + \mathbf{k}' - \mathbf{k}'' - \mathbf{k}''' - \mathbf{Q})
$$

=
$$
\int d^2 \mathbf{K} \delta(\mathbf{k} + \mathbf{k}' - \mathbf{K}) \delta[\mathbf{k}'' + \mathbf{k}''' - (\mathbf{K} - \mathbf{Q})],
$$
 (A3)

$$
\delta(\epsilon_{\mathbf{k}} + \epsilon_{\mathbf{k'}} - \epsilon_{\mathbf{k''}} - \epsilon_{\mathbf{k'''}})
$$

=
$$
\int dE \, \delta(\epsilon_{\mathbf{k}} + \epsilon_{\mathbf{k'}} - E) \delta(\epsilon_{\mathbf{k''}} + \epsilon_{\mathbf{k'''}} - E).
$$
 (A4)

Using the identity

$$
f_{\mathbf{k}}^{0} f_{\mathbf{k}'}^{0} (1 + f_{\mathbf{k}''}^{0}) (1 + f_{\mathbf{k}'''}^{0}) = e^{\beta (E - 2\mu)} f_{\mathbf{k}}^{0} f_{\mathbf{k}'}^{0} f_{\mathbf{k}''}^{0} f_{\mathbf{k}'''}^{0}, \tag{A5}
$$

we arrive at

$$
P = \frac{U^2 a^4}{16\hbar} \beta \int dE \, e^{\beta(E - 2\mu)} w(E),
$$

$$
w(E) = \int d^2 \mathbf{K} \, G_{\mathbf{K}}(E),
$$
 (A6)

$$
G_{\mathbf{K}} = \frac{1}{(2\pi)^5} \sum_{Q} \left[F_{\mathbf{K}}^{(2)} F_{\mathbf{K} - \mathbf{Q}}^{(0)} - 2F_{\mathbf{K}}^{(1)} F_{\mathbf{K} - \mathbf{Q}}^{(1)} + F_{\mathbf{K}}^{(0)} F_{\mathbf{K} - \mathbf{Q}}^{(2)} \right],
$$

$$
F_{\mathbf{K}}^{(m)}(E) = \int d^2 \mathbf{q} \left(\phi_{\frac{\mathbf{K}}{2} - \mathbf{q}} + \phi_{\frac{\mathbf{K}}{2} + \mathbf{q}} \right)^m f_{\frac{\mathbf{K}}{2} - \mathbf{q}}^0 f_{\frac{\mathbf{K}}{2} + \mathbf{q}}^0
$$

$$
\times \delta \left(\epsilon_{\frac{\mathbf{K}}{2} - \mathbf{q}} + \epsilon_{\frac{\mathbf{K}}{2} + \mathbf{q}} - E \right). \tag{A7}
$$

For each density and temperature we tabulate the 1D integral $F_{\mathbf{K}}^{(m)}(E)$ on a **K** and *E* grid. The remaining 3D integral in Eq. (A6) is then efficiently calculated as a discrete sum. We use a sequence of finer grids to verify that our result has converged to the continuum limit.

The chemical potential μ is fixed for each *n* and *T* by numerically solving

$$
n(\beta, \mu) = \int \frac{d^2 \mathbf{k}}{(2\pi)^2} f_{\mathbf{k}}^0,
$$
 (A8)

and the current *j* is expressed as

$$
j = -e \int \frac{d^2 \mathbf{k}}{(2\pi)^2} \phi_{\mathbf{k}} \frac{\partial f_k^0}{\partial \epsilon_k} (\mathbf{v}_{\mathbf{k}})_x
$$

=
$$
\frac{4t^2 a e}{\hbar} \beta \int \frac{d^2 \mathbf{k}}{(2\pi)^2} \sin^2(k_x a) e^{\beta(\epsilon_{\mathbf{k}} - \mu)} (f_{\mathbf{k}}^0)^2.
$$
 (A9)

One can interpret Eq. (A6) as a Boltzmann sum, where the weight function $w(E)$ corresponds to a density of scattering states. Figure $\overline{3}$ $\overline{3}$ $\overline{3}$ shows $w(E)$ for several temperatures, where *E* corresponds to the center-of-mass energy of the scattering particles. There are three features visible, a sharp peak at $E = 0$, a broader peak near $E = -4t$, and a diffuse background running from $-8t < E < 8t$. The $E = 0$ peak comes from the scattering between two particles that are both near the van Hove singularity of the square lattice. The $E = -4t$ peak is due to the scattering between a low-energy particle and one near the van Hove singularity. All other scattering events contribute to the background. As the temperature is lowered, and fewer particles reside near the van Hove singularity, the relative weight of these features shift. At low temperature the peak near $E = -4t$ dominates the calculation of *P*, giving rise to the exponential suppression of both ρ and Γ .

APPENDIX B: LOW-DENSITY LIMIT

Here we establish that the resistivity is well-defined in the low-density limit $n \to 0$, neither diverging nor vanishing. We first note that in this limit the chemical potential approaches $-\infty$, and hence, the Bose-Einstein distribution is well approximated by the Maxwell-Boltzmann expression

FIG. 3. The weight function $w(E)$ from Eq. [\(A6\)](#page-3-0) as a function of the energy *E* of the scattering particles.

 $f_{\mathbf{k}}^0 = (e^{\beta(\epsilon_{\mathbf{k}}-\mu)} - 1)^{-1} \approx e^{-\beta(\epsilon_{\mathbf{k}}-\mu)}$. Furthermore, we can remove the chemical potential dependence by considering the ratio

$$
\frac{f_{\mathbf{k}}^0}{n} = \frac{e^{-\beta \epsilon_{\mathbf{k}}}}{\int \frac{d^2 \mathbf{k}}{(2\pi)^2} e^{-\beta \epsilon_{\mathbf{k}}}}.
$$
(B1)

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Explicitly making this substitution, we write the resistivity as

$$
\rho = \frac{\frac{P}{n^4}}{\left(\frac{j}{n^2}\right)^2}.
$$
\n(B2)

As $n \to 0$, both the numerator and the denominator are independent of n , and hence, ρ approaches a constant.

APPENDIX C: VARIATIONAL PROCEDURE

According to Refs. [25,26], the variational parameter ξ is optimized by maximizing the entropy production from scattering, $T S_{\text{coll}} = \xi^2 P$ subject to the constraint $\xi X = \xi^2 P$, where $X = \mathbf{j} \cdot \mathbf{E}$. We therefore introduce a Lagrange multiplier λ and extremize

$$
\mathcal{L}[\xi] = \xi^2 P - \lambda(\xi^2 P - \xi X). \tag{C1}
$$

The conditions $\delta_{\xi} \mathcal{L} = 0$ and $\xi X = \xi^2 P$ give $\lambda = 2$, which allows us to write the Lagrangian in Onsager's form $\mathcal{L} =$ $-T\dot{S}_{\text{coll}} + 2T\dot{S}_{\text{field}}$ [30,31] and conclude that

$$
\xi = \frac{X}{P} = \frac{\mathbf{j} \cdot \mathbf{E}}{P}.
$$
 (C2)

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