Linear resistivity and Sachdev-Ye-Kitaev (SYK) spin liquid behavior in a quantum critical metal with spin-1/2 fermions

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"Strange metals" with resistivity depending linearly on temperature down to low T have been a long-standing puzzle in condensed matter physics. Here, we consider a lattice model of itinerant spin-1/2 fermions interacting via on-site Hubbard interaction and random infinite-ranged spin-spin interaction. We show that the quantum critical point associated with the melting of the spin-glass phase by charge fluctuations displays non-Fermi liquid behavior, with local spin dynamics identical to that of the Sachdev-Ye-Kitaev family of models. This extends the quantum spin liquid dynamics previously established in the large-M limit of SU(M) to SU(2) spin-1/2 electrons. Remarkably, the quantum critical regime also features a Planckian linear-T resistivity associated with a T-linear scattering rate and a frequency dependence of the electronic self-energy consistent with the marginal Fermi liquid phenomenology.

Planckian metals | strange metals | marginal Fermi liquid | Sachdev-Ye-Kitaev models | cuprate superconductors

**Significance**

In "Planckian metals," electrons dissipate energy at the fastest possible rate allowed by the fundamental laws of quantum mechanics, resulting in a linear temperature dependence of their electrical resistivity. Although observed for a number of quantum materials, this phenomenon lacks a general theoretical understanding and is often considered as one of the prominent fundamental questions in condensed matter physics. Here, we show that Planckian dissipation and a behavior consistent with the "marginal Fermi liquid" phenomenology emerge in the quantum critical regime separating a Mott insulating spin glass and a Fermi liquid. By establishing this behavior in an explicit model solvable by state-of-the-art computational methods, our theory paves the way toward a deeper understanding of Planckian or "strange" metals.

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which the SG phase can be quantum melted at the QCP and 2) providing an explicit numerical solution of this model directly for spin-1/2 SU(2) fermions. We find, remarkably, that the quantum critical regime displays SY spin-liquid correlations and a scattering rate linear in temperature, leading to T-linear resistivity down to $T = 0$ at the QCP. Our numerical results are consistent with the MFL phenomenology.

We consider a “t-U-J” model of itinerant spin-1/2, SU(2) fermions with an onsite repulsive-U Hubbard interaction and a random infinite-ranged spin–spin coupling, at half-filling. Using the extended dynamical mean-field theory framework (EDMFT) (29–32) and numerical methods detailed below, we obtain the phase diagram displayed in Fig. 1. At $t/U \rightarrow 0$, we have a Mott insulating SG phase (Fig. 1), where the fermions are localized onsite and the model reduces to the disordered Heisenberg model. SG order is found below a freezing temperature $T_g \approx 0.14J$ for $t/U = 0$ as previously established (28, 33) (SI Appendix, sections A and B). As $t/U$ is increased, the single-occupancy constraint is relaxed, and the charge fluctuations lead to quantum melting of SG order at a QCP ($t/U_c \approx 0.31$) separating the SG from an FL phase at low-enough temperature for ($t/U) > (t/U_c)$ (blue points in Fig. 1). Our key finding is a quantum critical region emanating from the QCP with QSL spin dynamics identical to that of the SY model (26) and T-linear MFL scattering rate $\text{Im}\Sigma(\omega \rightarrow 0, T) \propto T$ (red points in Fig. 1), leading to $T$-linear resistivity as shown below.

More precisely, our model Hamiltonian reads

$$H = -\sum_{\langle ij \rangle, s=\uparrow, \downarrow} t_{ij} c_{is} c_{js} + U \sum_i n_{i\uparrow} n_{i\downarrow} - \sum_{i<j} J_{ij} \vec{S}_i \cdot \vec{S}_j,$$  \[1\]

In this expression, $J_{ij}$ are quenched random Heisenberg interactions (33) drawn from a Gaussian distribution with $\langle J_{ij} \rangle = 0$ and $\left \langle J_{ij}^2 \right \rangle = J^2$, $N'$ is the number of sites, and $\vec{S}_i = \frac{1}{2} c_{is}^{\dagger} \vec{\sigma} s^s c_{is}$, with $\vec{\sigma}$ the Pauli matrices. The model can be formulated either on the infinite connectivity $z \rightarrow \infty$ Bethe lattice with $t_0 = t/\sqrt{z}$ or on a fully connected lattice with Gaussian distributed random $t_0$ with $\langle t_0 \rangle = 0$ and $\langle t_0^2 \rangle = t^2/\mathcal{N}$, leading to identical equations in the phase without magnetic ordering after replica averaging (34). We restrict ourselves to the half-filling case $\mu = U/2$ and choose $J = 0.5t$. A study of the $SU(M)$ version of this model in the large-$M$ limit at half-filling is in ref. 35.

To investigate the phase diagram of model [1], both the onsite repulsion in the charge channel and the random interaction in the spin channel need to be tamed. This is achieved using the EDMFT framework and the replica trick. In this framework, the calculation of the local Green’s function and spin–spin correlation function is mapped onto the solution of a local “quantum impurity” problem subject to a self-consistency condition (26, 27, 29, 33, 34, 36–38). This mapping is exact in the infinite connectivity $z \rightarrow \infty$ or infinite volume limit $\mathcal{N} \rightarrow \infty$ of the two formulations of the model discussed above.

The resulting local effective action, after disorder averaging and making a replica diagonal ansatz, reads

$$S_{\text{eff}} = -\beta \sum_{i, \alpha} c_{i\alpha}^{\dagger} (i\omega_n + \mu - \Delta(i\omega_n)) c_{i\alpha} + U \int_0^\beta d\tau \sum_i \bar{n}_i,$$

$$- \frac{J^2}{2} \int_0^\beta d\tau d\tau' \text{Q}(\tau - \tau') \langle \vec{S}(\tau) \cdot \vec{S}(\tau') \rangle.$$  \[2\]

In this expression, $\beta = 1/T (k_B = 1)$ is the inverse temperature, $\tau \in [0, \beta]$ stands for imaginary time, and $\omega_n = (2n + 1)\pi/\beta$ are Matsubara frequencies. The dynamical mean-field (hybridization) function $\Delta$ and effective spin–spin retarded interaction $Q$ are subject to the following self-consistency conditions:

$$\Delta(\tau) = t^4 G(\tau), \quad \text{Q}(\tau - \tau') = \frac{1}{3} \left \langle \hat{S}_s(\tau) \cdot \hat{S}_s(\tau') \right \rangle,$$  \[3\]

in which the local Green’s function $G(\tau) \equiv -\langle T_c_{\alpha}(\tau) c_{\alpha}^{\dagger}(0) \rangle$ and the local spin–spin correlator $\left \langle \hat{S}_s(\tau) \cdot \hat{S}_s(\tau') \right \rangle$ are to be computed with the local effective action [2]. Noting that $i\omega_n + \mu - \Delta(i\omega_n)$ is the inverse effective one-body propagator of this action, a fermionic self-energy can be defined from Dyson’s equation as

$$\Sigma(i\omega_n) = i\omega_n + \mu - \Delta(i\omega_n) - \frac{1}{\text{Q}} \left \langle \hat{S}(\tau) \cdot \hat{S}(\tau) \right \rangle.$$  \[4\]

The local action [2] still presents a strongly correlated problem. SY (26) made further progress on the random Heisenberg model by extending the spin symmetry to $SU(M)$ and taking the $M \rightarrow \infty$ limit, which allows for an analytical calculation of the spin–spin correlator of [2] and reduces the self-consistent problem to a nonlinear integral equation. This $M \rightarrow \infty$ limit was extended to itinerant fermions within the t-J$_ij$ model by Parcollet and Georges (PG) (27), who obtained an FL regime of the doped model at low $T$, and a quantum critical regime associated with the proximity of the spin-liquid Mott insulator characterized by a $\sqrt{\omega} - \sqrt{T}$ self-energy but remarkably, bad metal behavior with linear resistivity. Recently, fermionic versions of the random coupling problem, the so-called Sachdev-Ye-Kitaev (SYK) models (39, 40), garnered much interest with again a soluble limit for a large number of flavors $M \rightarrow \infty$.

Recent works (41–45) extended the mechanism of PG (27) for linear-$T$ resistivity to a lattice of SYK “quantum dots” with hopping. Interestingly, when SYK dots are coupled to another band of otherwise free and translationally invariant (uniform hopping) fermions, not only does the $T$-linear resistivity extend down to zero temperature, but the mechanism switches to that driven by the MFL T-linear scattering rate (46, 47).

For the physical limit of a single flavor of spin-1/2 fermions that is of our interest, the self-consistency equations above require computing two- and four-point correlators in the local

![Fig. 1. Calculated phase diagram of the t-U-J model [1] at $J/t = 0.5$. Solid black curve indicates a second-order phase transition to SG order. Round markers represent parameters for which we have explicitly solved the model. Markers have been colored red where we find a quantum critical metal (QCM) with QSL spin dynamics and blue where we find an FL. Background shading interpolates between the explicitly solved points. Dashed black curve indicates the cross-over between QCM and FL regimes. Gray markers indicate Mott insulating solutions. Black markers indicate SG ordered solutions.](18342.png)
model with \( SU(2) \) symmetry. We use an implementation of Rubtsov's continuous-time interaction-expansion quantum Monte Carlo (CT-INT) (48) algorithm, which is based on the TRIQS library (49). The algorithm works in imaginary time, so we will discuss most of our results directly on the imaginary axis without analytic continuation, except in the discussion of transport. Our implementation determines the local spin–spin correlator from the impurity three-point vertex function rather than through an operator insertion measurement. This algorithmic improvement allows for a drastic speedup of the calculations.

Let us first consider the long-time spin dynamics. In Fig. 2, we display the local spin–spin correlation function \( Q(\tau) \) at a fixed low temperature \( T/t = 0.01 \) for various \( t/U \) approaching the QCP at \( (t/U)_c \approx 0.31 \) from the FL limit cutting the phase diagram Fig. 1 along the horizontal axis. In Fig. 2, Inset, we also display how \( Q(\tau) \) varies upon raising temperatures for fixed \( t/U = 0.357 \) making a vertical cut in the phase diagram slightly away from the QCP. Since we work in the Matsubara formalism, a zero-temperature long-time asymptotic form \( Q(t) \sim 1/t^\alpha \) transforms into a scaling function \( Q(t) \sim (\pi/\beta) \sin(\pi/\beta) \alpha \), and the data should be examined away from \( \tau = 0, \beta \). Examining correlators at \( \tau = \beta \) is an established method of analyzing long-time, low-frequency behavior (50). However, as we are interested in the scaling exponent, we examine the full \( \tau \) dependence of the spin–spin correlator in a large region around \( \tau = \beta/2 \). Away from the critical point, for \( t/U > 1.0 \), we obtain the FL behavior at long time \( Q(t) \sim 1/t^\alpha \) \((\alpha = 2)\). The closer one gets to the critical point, the longer it takes to reach this asymptotic regime, reflecting the decrease of the FL coherence scale close to the critical point. Once in the quantum critical regime, for \( t/U \approx (t/U)_c \approx 0.31 \), the long-time spin dynamics crosses over to \( Q(t) \sim 1/t \) \((\alpha = 1)\), which is the same power law as in the SY \( M = \infty \) model. The QSL to FL cross-over is also visible in the temperature cut shown in Fig. 2. Inset, where we observe the cross-over from \( 1/t \) within the quantum critical fan above the FL coherence temperature to \( 1/t^\alpha \) at low temperatures. The phase classification at each point in Fig. 1 follows the above criterion to identify the FL regime and the QSL regime.

These results establish that our \( SU(2) \) \( t-U-J \) model has, in the quantum critical regime, the same QSL local spin dynamics \((\alpha = 1)\) as the SY model in the \( M = \infty \) limit. Renormalization-group (RG) methods should prove useful in establishing analytically our numerical findings for \( SU(2) \). For simplified versions of the effective action [2] (e.g., involving only localized spins) (51), RG methods have indeed established (51–61) that the \( Q(t) \sim 1/t \) spin-liquid behavior is the only one consistent with the self-consistency condition [3]. This was recently extended to the QCP obtained by doping the \( U = \infty \) model (62).

Let us now consider the one-particle properties, encoded by the self-energy \( \Sigma \). In the FL regime for \((t/U)_c \ll (t/U)_c \), the self-energy has the low-energy expansion:

\[
\Sigma(\omega_n, T) \approx \left(1 - \frac{1}{Z}\right) \omega_n + \frac{\omega_n^2 - (\pi/\beta)^2}{E} + O(\omega_n^3). \tag{5}
\]

In the small hopping limit \((t/U) \ll (t/U)_c\), \( \Sigma \) diverges at low frequencies as \( 1/\omega_n \), indicating a transition into an insulating phase (SI Appendix, section C). We examine the cross-over from the FL to the quantum critical regime in several ways. First, a direct consequence of [5] is that the self-energy at the first Matsubara frequency is linear in temperature with vanishing quadratic corrections (63): \( \text{Im} \Sigma(\omega_0 = i\pi/\beta) \approx (1 - 1/\pi T)E + O(\tau^2) \). Deviation from linearity in temperature \( T/t \) signals the FL coherence scale and hence, the cross-over to the quantum critical regime. This is illustrated in Fig. 3A, where \( t/U \) approaches \((t/U)_c\), the self-energy increases, and \( T/t \) (indicated by arrows on the figure) decreases. More precisely, we extract the quasiparticle residue \( Z \) and the coherence scale \( E \) by fitting the functional form [5] to the low-energy data using weighted least squares. Fig. 3B shows that \( Z \) and \( E \) vanish at the QCP. The susceptibility to SG order is given by \((37, 38)\) \( \chi_{sg} \propto \chi^2(1 - J^2x^2) \) with \( \chi \) the local susceptibility. As shown in Fig. 3, we find that \( 1 - Jx \) also vanishes close to the QCP, indicating the boundary of the SG phase. Within our numerical accuracy, we cannot however exclude that \( 1 - Jx \) vanishes at a slightly larger value of \( t/U \) than \( E \), possibly indicating a small intervening region of metallic SG (62).

In order to analyze the OCP, we attempt to scale the self-energy for \((t/U)_c \) close to \((t/U)_c\), for our lowest temperature \( T/t = 0.01 \), with an ansatz of the form

\[
\text{Im} \Sigma(\omega_n) \approx \text{Im} \Sigma(0) + f \left(\frac{\omega_n}{\omega_*}\right), \tag{6}
\]

which applies for \( \omega_n \) and \( \omega_* \) smaller than the high-energy cutoff \( J \) but \( \omega_n/\omega_* \) otherwise arbitrary. We determine numerically \( \text{Im} \Sigma(0) \), \( \omega_* \), and the scaling function \( f \) by requesting that optimal data collapse is obtained, using a minimization procedure. We obtain a remarkable collapse of the data, presented in Fig. 4A, with \( \omega_* \) presented in Fig. 3B.

For \( \omega < \omega_* \), the ansatz [6] has to reproduce [5], which implies \( Z \propto \omega \) for \( \omega \rightarrow 0 \), and \( E \propto \omega^2 \), hence, \( E \propto Z^2 \), as illustrated in Fig. 3B, Inset. Note however that the \( \omega_* \) obtained from the data collapse does not perfectly vanish close to the OCP, which may be due to numerical uncertainty, or possibly to a weakly first-order transition or to an intervening metallic SG phase as mentioned above. In the quantum critical regime (i.e., for \( \omega > \omega_* \)), the self-energy is very well described by an MFL form \( \text{Im} \Sigma(\omega_n) \propto \Sigma(0) + a_{\omega_n} \text{Im} \text{Im} \Sigma(0) \) (Fig. 4A, Inset). However, the low-temperature behavior obtained in the large-\( M \) limit (26) (i.e., \( \sqrt{\omega_n} \) cannot be excluded given our data. Indeed, the CT-INT algorithm is faced with a sign problem at low \( T \), which prevents us from reaching the very low-temperature regime.
required to settle this question. This conclusion holds both for the scaling function $f$ and for a direct analysis of the self-energy at $t/U = (t/U)_{\infty}$.

The value of the self-energy at zero-frequency Im$\Sigma(0)$ is of crucial importance for transport properties. In Fig. 4C, we show Im$\Sigma(0)$ extracted from the scaling analysis, for various $U$ close to the QCP. We find that Im$\Sigma(0) \sim T$ at low temperature at the QCP. This is confirmed in Fig. 4D: Im$\Sigma(\omega)$ obtained (by interpolation) for fixed imaginary frequency $\omega$ is linear with temperature, with a slope weakly dependent on the frequency.

Let us finally turn to the direct current (DC) resistivity in the quantum critical region. The Kubo formula reduces to the polarization bubble (vertex corrections vanish in this quantity in Dynamical Mean Field Theory [DMFT]):

$$\sigma_{DC} = \frac{2\pi e^2}{h} \int d\omega \frac{\beta}{4 \cosh^2(\beta \omega/2)} \int d\epsilon \phi(\epsilon) A(\epsilon, \omega)^2. \quad [7]$$

This expression only relies on vanishing vertex corrections, which is valid in DMFT as well as in other contexts, such as large-$N$ SYK models. In this expression, $\epsilon$ is the energy of a bare single-particle state within the band, $A(\epsilon, \omega) = -(1/\pi)\text{Im} G(\epsilon, \omega)$ is the energy-(momentum-)resolved spectral function, and $\phi(\epsilon)$ is the transport function $\phi(\epsilon) = \sum_k \frac{\partial q_k}{\partial \epsilon_k} \delta(\epsilon - \epsilon_k)$, which we take to be the sum-rule preserving expression on the Bethe lattice (e.g., ref. 10): $\phi(\epsilon) = \phi(0)[(1 - (\epsilon/2t)^2]^{1/2}$. To obtain $\sigma_{DC}$, we perform an analytic continuation of the Monte Carlo data using Padé approximants (64) to obtain the real-frequency self-energy $\Sigma(\omega) = \Sigma^R(\omega) + i\Sigma^I(\omega)$ and the spectral function: $\pi A(\epsilon, \omega) = -\Sigma'(\omega)/[(\omega + \mu - \epsilon - \Sigma^R(\omega))^2 + \Sigma^I(\omega)^2]$. The resulting resistivity $\rho_{DC} = 1/\sigma_{DC}$ vs. temperature $T$ is plotted in Fig. 4C, clearly consistent with $T$-linear resistivity within numerical accuracy.

The origin of this behavior can be directly related to the $T$-linear behavior of the scattering rate $\Sigma'(0)$. Indeed, observing that the latter is a much smaller scale than the bandwidth at low $T$, the integral over $\epsilon$ can be approximated as

$$\int d\epsilon \phi(\epsilon) A(\epsilon, \omega)^2 \sim \frac{\phi(\omega + \mu - \Sigma^R(\omega))}{2\pi i \Sigma^I(\omega)}.$$ \quad [8]

Due to the Fermi factor only, $|\omega| \lesssim T$ is relevant for the frequency integral, so that the right-hand side of this expression can be replaced by its Fermi surface contribution $\omega = 0$ (SI Appendix, section D). Observing that $\mu - \Sigma^R(0) = 0$, we finally obtain

$$\sigma_{DC} = \frac{\epsilon^2 \phi(0)}{h} \frac{\beta \int d\omega}{\int (\beta \omega/2) \Sigma^I(\omega)} \sim \frac{\epsilon^2 \phi(0)}{h T}. \quad [9]$$

$\rho_0 = (h/2\pi)^2/\phi(0)/t$ can be taken as the order of magnitude of the MIR resistivity (10), so that we obtain at the QCP $\rho_{DC}/\rho_0 \sim T/t$ down to the lowest value of $T$ we could reach.

We would like to emphasize that both the mechanism and the physical meaning of this $T$-linear resistivity are different from the ones reported in ref. 27 and in the SYK $M \to \infty$ lattice models (41–45). There, the scattering rate had a $\sim \sqrt{T}$ temperature dependence and dominated the band dispersion in the incoherent metal regime $T > T^*$, resulting in the resistivity being proportional by the square of the scattering rate and larger than the MIR value. Here, in contrast, the scattering rate is $T$ linear (Planckian) and small at low $T$, and the band dispersion dominates, resulting in linear resistivity down to low $T$. The present mechanism is also distinct from the generic bad metal behavior of lattice models at very high $T$ comparable with the bandwidth (7, 8, 10, 11, 14): there, the scattering is constant, and the $T$-linear behavior is associated with the $T$ dependence of thermodynamic quantities such as the kinetic energy $\sim 1/T$, which play the role of an effective carrier number. We have checked (SI Appendix, section E) that in contrast the kinetic energy of our model is constant in the range of $T$ of interest.

In this work, we considered the insulator to metal transition and quantum melting by charge fluctuations of the SG ground state of the $SU(2)$ random-bond Heisenberg model. At the QCP separating the SG from the FL, we find a non-FL state with long-wavelength spin correlations $\langle \hat{S}(\cdot) \cdot \hat{S}(0)\rangle \sim 1/t$ [as in the large-$M$ limit of the $SU(M)$ SY model] and a $T$-linear resistivity arising from a $T$-linear (Planckian) scattering rate $\text{Im} \Sigma(\omega = 0, T) \propto T$. In the temperature range accessible in this work, this quantum critical regime is compatible with a marginal Fermi phenomenology $\Sigma(\omega) \sim \omega \log \omega$. The Planckian scattering rate and MFL behavior are driven by local quantum critical fluctuations from the
QCP. Fully establishing this behavior down to zero temperature may require a new generation of quantum impurity solvers, such as real-time diagrammatic Monte Carlo (65, 66). Another open question is whether our results for the scattering rate also apply to the doped case recently considered in ref. 62. Also, finding the RG fixed point associated with our metal insulator transition QCP remains an open question.

Data Availability. The data analyzed in this article are available at the Figshare repository (67).

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Fig. 4. (A) Imaginary part of self-energy with the scattering rate subtracted –(ImΣ(ω)/ω) – (ImΣ(0)) vs. the scaled frequency ω/ω* for various values of T/U near the QCP at T/t = 0.01, demonstrating the collapse onto the universal scaling function f(ω/ω*) (gray solid curve). Color scheme follows B. (Inset) Imaginary part of self-energy –(ImΣ(ω)) vs. Matsubara frequencies ω_q at the QCP T/U = 0.312 and lowest accessible temperature T/t = 0.01. Also shown are low-frequency fits of self-energy to the MFL form ω_c + dω_c log ω_q/b (orange) and the SYK form ω_c + a_0 + b_0ω_q (green). (B) Scattering rate –(ImΣ(0))/T vs. temperature T/t at various values of T/U near the QCP. At the QCP (T/U = 0.312, green), the scattering rate is T linear (linear fit in gray), in contrast to the quadratic behavior in the FL regime (blue). (C) Resitivity ρ_DC/ρ0 vs. temperature T/t at the QCP computed with the analytically continued Green’s function. The unit of resistivity is the MIR value ρ_0 = e^2/h, where e is the transport function. (D) Imaginary part of self-energy at fixed, interpolated values of Matsubara frequency –(ImΣ(ω) = fixed, T) vs. temperature T/t at the QCP T/U = 0.312, for various fixed values of frequency.

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