

# Linear resistivity from hydrodynamics

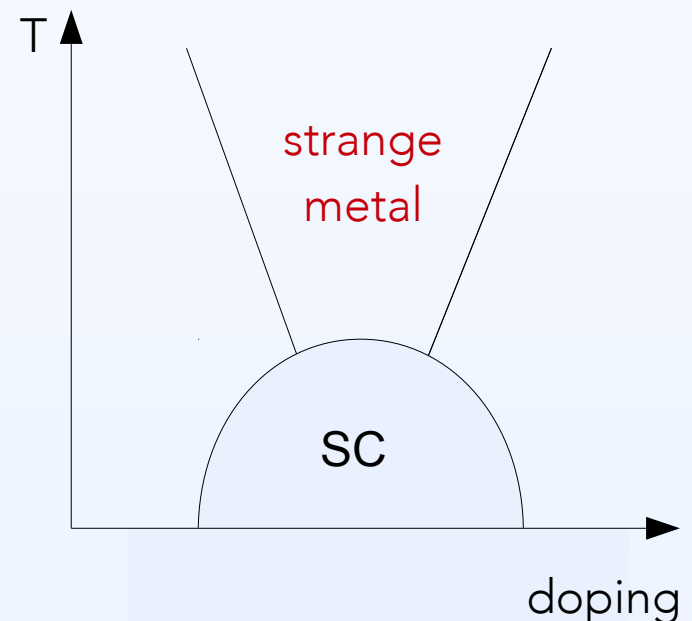
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Based on: 1311.2451 [hep-th] by RD, K. Schalm, J. Zaanen

# Linear resistivity in the cuprates

- The **strange metal** state of the high- $T_c$  cuprate superconductors has weird transport properties.
- The most famous is that its **resistivity is linear in temperature**.
- Why? There is a non-trivial IR fixed point. It is not a Fermi liquid. **What is it?**
- Taking inspiration from holography, I will describe a very simple mechanism which produces a resistivity like this.



# Linear resistivity from holography

- Consider the classical theory of gravity with action

$$S = \frac{1}{2\kappa_4^2} \int d^4x \sqrt{-g} \left( R - \frac{1}{4} e^\phi F_{\mu\nu} F^{\mu\nu} - \frac{3}{2} \partial_\mu \phi \partial^\mu \phi + \frac{6}{L^2} \cosh \phi \right)$$

see e.g. Gubser, Rocha 0911.2898

- This has a charged black brane solution which can be uplifted to a solution of 11D supergravity. Like the cuprate strange metals, it has an entropy linear in  $T$ .
- Introducing a **random distribution of impurities or a periodic lattice** in this state produces a resistivity which is approximately linear in  $T$ .

Anantua, Hartnoll, Martin, Ramirez (2012)

# How can this be realistic?

- How can they possibly be related? A priori, this field theory looks totally unrelated to the cuprates.
- The mechanism which produces a linear resistivity is **independent of many details of the field theory**.
- **It does not require holography**. It can be understood from general principles of strongly interacting quantum critical states.
- The holographic state is just an example of where this mechanism is at work. This is not so dissimilar to the role of holography in understanding the QGP.

# Outline of the talk

- Resistivity in states with an almost conserved momentum
- Momentum dissipation rate from dynamics near black brane horizon
- Momentum dissipation rate from hydrodynamics
- Linear resistivity from hydrodynamics

# Slow momentum dissipation I

- DC transport properties, like the resistivity, tell us about the late time response of a system to an external source.
- In theories where long-lived quasiparticles carry the current, the quasiparticle decay rate controls the resistivity.
- If there are no long-lived quasiparticles (e.g. in a strongly interacting quantum critical theory), the current intrinsically wants to decay quickly.
- But in a system with perfect translational invariance, momentum is conserved. If the current carries momentum, it cannot decay.  
Therefore  $\rho_{DC} = 0$

# Slow momentum dissipation II

- Suppose, in a system like this, translational invariance is broken in a weak way so that **momentum dissipates slowly**.
- This will cause the current to decay slowly at a rate controlled by the momentum dissipation rate  $\rho_{DC} \sim \Gamma$
- As translational invariance is broken weakly, the momentum dissipation rate  $\Gamma$  can be calculated **perturbatively**.
- Suppose we turn on a lattice i.e. a spatially periodic source for an operator in the IR

$$\delta H = V \int dx e^{ik_L x} \mathcal{O}(x)$$

# Slow momentum dissipation III

- At leading order, the rate at which momentum dissipates into the lattice is determined by the spectral weight in the translationally invariant system

$$\Gamma = \frac{V^2 k_L^2}{\chi_{\vec{P}\vec{P}}} \lim_{\omega \rightarrow 0} \frac{\text{Im} G_{\mathcal{O}\mathcal{O}}(\omega, k_L)}{\omega} \Big|_{V \rightarrow 0} \quad \text{Hartnoll, Hofman (2012)}$$

- This tells us the number of low energy degrees of freedom of the system at the lattice momentum  $k_L$ . It is these that will couple to the lattice, once it is turned on.
- If a spatially random source for an operator is turned on,

$$\Gamma = \frac{\bar{V}^2}{\chi_{\vec{P}\vec{P}}} \int d^2 k \, k^2 \lim_{\omega \rightarrow 0} \frac{\text{Im} G_{\mathcal{O}\mathcal{O}}(\omega, k)}{\omega} \Big|_{\bar{V} \rightarrow 0} \quad \begin{array}{l} \text{Hartnoll et. al. (2007)} \\ \text{Hartnoll, Herzog (2008)} \end{array}$$



# Slow momentum dissipation: summary

- If we have a charged state in which the only long-lived quantity is the momentum, the **resistivity is proportional to the momentum dissipation rate**.
- At leading order, this is determined by properties of the translationally invariant state.
- Although this is independent of holography, it is applicable to some of the field theory states described by holography.
- In these cases, we can use holography to calculate the response functions that control the momentum dissipation rate and resistivity.

# Holography: gravitational solution

- Using these tools, it was found that the state dual to the charged black brane solution to the Einstein-Maxwell-Dilaton theory

$$S = \frac{1}{2\kappa_4^2} \int d^4x \sqrt{-g} \left( R - \frac{1}{4} e^\phi F_{\mu\nu} F^{\mu\nu} - \frac{3}{2} \partial_\mu \phi \partial^\mu \phi + \frac{6}{L^2} \cosh \phi \right)$$

has  $\rho_{DC} \sim T$  when coupled to periodic, or spatially random, sources of charge density or energy density.

Anantua, Hartnoll, Martin, Ramirez (2012)

- The relevant gravitational solution is

$$ds^2 = \frac{r^2 g(r)}{L^2} (-h(r) dt^2 + dx^2 + dy^2) + \frac{L^2}{r^2 g(r) h(r)} dr^2, \quad A_t(r) = \mu \left( 1 - \frac{Q + r_0}{Q + r} \right),$$

$$\phi(r) = \frac{1}{3} \log(g(r)), \quad g(r) = \left( 1 + \frac{Q}{r} \right)^{\frac{3}{2}}, \quad h(r) = 1 - \frac{(Q + r_0)^3}{(Q + r)^3},$$

# IR geometry of charged black brane

- The near horizon geometry is conformal to  $\text{AdS}_2 \times \mathbb{R}^2$ . In the usual classification of near-horizon geometries, it has

$$z \rightarrow \infty \quad \text{with} \quad \theta/z = -1$$

- It is similar to the near-horizon  $\text{AdS}_2 \times \mathbb{R}^2$  geometry of RN- $\text{AdS}_4$ . The main difference is that this state has entropy

$$s \sim T$$

- $z \rightarrow \infty$  means **local quantum criticality** in the field theory: the low energy physics is approximately momentum-independent.
- Greens functions of fields in this IR geometry have the generic form
$$\lim_{\omega \rightarrow 0} \text{Im} \mathcal{G}^{IR}(\omega, k) \sim \omega T^{2\nu(k)-1}$$

# Spectral functions from gravity

- Linear perturbations of the energy density  $T^{tt}$  and charge density  $J^t$  are irrelevant in the IR: spatially periodic or random sources will cause **momentum to dissipate slowly**.
- The Greens functions can, in principle, be obtained from a matching calculation

$$G(\omega, k) = \frac{A(k) + B(k)\mathcal{G}^{IR}(\omega, k) + \dots}{C(k) + D(k)\mathcal{G}^{IR}(\omega, k) + \dots}$$

- The matching does not have to be done explicitly. At low T, the leading dissipative term is proportional to the IR Greens function

$$\text{Im}G(\omega \rightarrow 0, k) \sim H(k) \text{Im}\mathcal{G}^{IR}(\omega, k) \sim \omega T^{2\nu(k)-1} H(k)$$

# Linear resistivity from disorder

- The momentum dissipation rate due to neutral or charged disorder is:

$$\Gamma \sim \int d^2k \, k^2 \lim_{\omega \rightarrow 0} \frac{\text{Im}G(\omega, k)}{\omega} \sim \int d^2k \, k^2 H(k) T^{2\nu(k)-1} \sim T^{2\nu(0)-1}$$

- The homogeneous ( $k=0$ ) mode dominates the integral at low temperatures.

Anantua, Hartnoll, Martin, Ramirez (2012)

- This gives a DC resistivity  $\rho_{DC} \sim T$  because an analysis of mass terms in the near horizon geometry shows that the scaling dimension of  $T^{tt}$  and  $J^t$  is

$$\nu(k) = \sqrt{\frac{11}{3} + 2\hat{k}^2} - \frac{8}{3}\sqrt{1 + \frac{3}{2}\hat{k}^2} = 1 + O(\hat{k}^4)$$

- Finite momentum contributions to  $\nu(k)$  are small and give logarithmic corrections to  $\rho_{DC}$ :  $\rho_{DC}(T) \sim T / \log(\mu/T)$

# Linear resistivity from a lattice

- The momentum dissipation rate due to a neutral or charged lattice is:

$$\Gamma \sim k_L^2 \lim_{\omega \rightarrow 0} \frac{\text{Im}G(\omega, k_L)}{\omega} \sim k_L^2 H(k_L) T^{2\nu(k_L)-1} \sim T^{2\nu(k_L)-1}$$

- Provided the lattice momentum is of the order of the chemical potential (or less), there is an approximately linear DC resistivity

$$\rho_{DC} \sim T \quad \text{Anantua, Hartnoll, Martin, Ramirez (2012)}$$

- Again, it is because the finite  $k$  corrections to the dimension are small e.g.  $\nu(\hat{k} = 0.5) \sim 1.02$

$$\nu(\hat{k} = 1) \sim 1.2$$

# Brief summary of these results

- Without reference to holography, we can summarise why this state has a linear resistivity:
- A lattice or random disorder causes momentum to dissipate slowly.
- The dissipation rate is determined by the two-point functions of  $T^{tt}$  and  $J^t$  in the translationally invariant, locally critical state.
- At low  $T$ , these are approximately proportional to  $T$  because  $T^{tt}$  and  $J^t$  have dimension  $\nu(k) = 1 + O(k^4)$ .
- Generally, one finds power laws for locally critical states  $\rho_{DC} \sim T^\eta$

# A different perspective

- Why do these correlators have a term which is approximately linear in  $T$ ??? There is another way to understand it.  
RD, Schalm, Zaanen, 1311.2451
- We have learned a lot about the general principles of how charge and momentum are transported in holographic theories with translational invariance.
- These general principles appear to be true in real strongly interacting systems: they do not require the existence of a dual classical gravity description.
- This highlights a simple mechanism that can produce linear resistivity and which may be at work in real systems.



# Some history

- The simplest case: a black brane dual to a neutral, thermal state.
- At long distances and low energies  $\omega, k \ll T$ , these behave like hydrodynamic fluids with a minimal viscosity

$$\eta = \frac{\hbar}{k_B} \frac{s}{4\pi}$$

Kovtun, Son, Starinets (2004)  
Iqbal, Liu (2008)

- A small viscosity means that a fluid thermalises very quickly.  
e.g. in a kinetic theory of quasiparticles,  $\eta \sim$  mean free time
- It is not so surprising that a state with a holographic dual forms a hydrodynamic state in a short time.

# Hydrodynamics

- Hydrodynamics is an **effective theory**, telling us what the collective properties of the system are at long distances and low energies.

- For a relativistic fluid with  $\epsilon = 2P$ ,

$$T^{\mu\nu} = (\epsilon + P) u^\mu u^\nu + P g^{\mu\nu} - \eta \Delta^{\mu\alpha} \Delta^{\nu\beta} (\partial_\alpha u_\beta + \partial_\beta u_\alpha - g_{\alpha\beta} \partial_\gamma u^\gamma) + \dots$$

$$J^\mu = \rho u^\mu - \sigma_Q T \Delta^{\mu\nu} \partial_\nu (\mu/T) + \dots$$

- At leading order in spatial derivatives, **dissipation** is controlled by two transport coefficients: shear viscosity  $\eta$  and “universal conductivity”  $\sigma_Q$ .
- Their values depend upon the specific microscopic theory

# Greens functions from hydrodynamics

- These hydrodynamic equations tell us how the state will respond to small perturbations.
- They fix the form of the Greens functions at long distances and low energies e.g.

$$G_{\vec{P}_\perp \vec{P}_\perp}(\omega, k) = \frac{\eta k^2 + \dots}{i\omega - Dk^2 + \dots} \qquad D = \frac{\eta}{\epsilon + P}$$

- The shear viscosity controls the rate at which momentum diffuses and the universal conductivity controls the rate at which charge diffuses.

# Hydrodynamics of locally critical states I

- At long distances and low energies, hydrodynamics is a good **approximate** description of locally critical holographic states.
- Greens functions can be calculated by matching the IR Greens functions to the asymptotically AdS UV region.

$$G(\omega, k) = \frac{A(k) + B(k)\mathcal{G}^{IR}(\omega, k) + \dots}{C(k) + D(k)\mathcal{G}^{IR}(\omega, k) + \dots}$$

- This can be done numerically or, in some cases, analytically.
- Unlike the neutral case, hydrodynamics is a good approximate description even at low temperatures, provided that  $\omega, k \ll \mu$

see e.g. Edalati, Jottar, Leigh (2010), RD, Parnachev (2013)  
Tarrio (2013) and others

# Hydrodynamics of locally critical states II

- In the simplest case of RN-AdS, the matching can be done explicitly and analytically for some operators.

RD, Parnachev (2013)

- Ignoring finite  $k$  corrections to  $\nu(k)$  in the IR geometry, the correlation functions are just those of hydrodynamics, with certain values of the transport coefficients.
- These corrections are not important for the leading order resistivity in the presence of disorder or a lattice.
- The key point is that if a theory obeys hydrodynamics, the IR dimensions of operators are not random numbers: they are related to the transport coefficients.

# Hydrodynamics of locally critical states III

- The T dependence of Greens functions in a hydro theory are controlled by the T dependence of the transport coefficients.
- We have replaced one aspect of microscopic physics (operator dimensions) with another: values of transport coefficients.
- This is a complimentary view of the same situation.
- It is advantageous for one reason: we can make an informed estimate of the size of one of these transport coefficients in general

$$\eta \sim \frac{\hbar}{k_B} s$$

# Viscous contribution to resistivity

- There are many hydrodynamic contributions to the resistivity which will depend upon microscopic details of the theory.

neutral  $\rho_{DC} \sim \frac{\mathcal{V}_{Ttt}^2}{\sigma^2} \int dk k (\eta k^2 + \dots),$

charged  $\rho_{DC} \sim \frac{\mathcal{V}_{Jt}^2}{\sigma^2} \int dk k \left( \frac{1}{\sigma_Q} \left[ 2 \frac{\sigma^2}{\epsilon + P} - \left( \frac{d\sigma}{d\mu} \right)_T \right]^2 + k^2 \frac{\sigma^2}{(\epsilon + P)^2} \eta + \dots \right)$

+ analogous expressions for lattice deformations

- We will concentrate on the **viscous** term.
- A simple argument of why it exists is that momentum diffuses in a hydrodynamic liquid with diffusion constant  $D = \eta / m_e n_e$ .
- If translational invariance is broken over a length scale  $l$ , the time it takes for the momentum to dissipate is  $\tau^{-1} = D / l^2$

# Resistivity = entropy

- The memory matrix calculation confirms this. It has also been observed by other methods e.g.

$$\rho_{2D} = \frac{1}{2e^2} \left\langle \frac{T}{\kappa} (\delta s_0)^2 + (\eta + \zeta) \left( \nabla \frac{1}{n_0} \right)^2 \right\rangle.$$

From 1011.3068 [cond-mat.mes-hall]  
by A. Andreev, S. Kivelson, B. Spivak

- If a theory behaves like a **hydrodynamic liquid with minimal viscosity** down to the length scale over which impurities/the lattice are present, it will have a viscous contribution to its resistivity

$$\rho_{DC}(T) \sim \eta(T) \sim s(T)$$

provided that momentum is almost conserved.

- The locally critical states of holography obey this “entropy law”.



# Fermi liquids

- Why do conventional metals not have  $\rho_{DC}(T) \sim s(T)$ ?
- These do not behave hydrodynamically at long times. The quasiparticle interaction rate is small:  $\Gamma_{ee} \sim T^2$
- The corresponding viscosity is large:  $\eta_{FL} \sim \frac{1}{T^2}$
- This means it takes a long time  $\tau \sim T^{-2}$  for a Fermi liquid to equilibrate via interactions and form a hydrodynamic state.
- The electrons lose their momentum via interactions with the ionic lattice before the hydrodynamic state forms.

# Cuprates

- Strong electronic interactions cause the formation of a hydrodynamic state with a minimal viscosity over a short time scale. This hydro description applies at distances  $\sim \mu^{-1}$ .
- Slow momentum-dissipating interactions then produce a resistivity
$$\rho_{DC}(T) = \frac{A\hbar}{\omega_p^2 m_e l^2} \frac{S(T)}{k_B} \sim T$$
- This requires a **small** length scale  $\sim l \sim 10^{-9}\text{m}$
- **But** there is no residual ( $T=0$ ) resistivity as, in this limit, the electrons behave as a perfect fluid.
- This is radically different from FL theory: **it should be testable.**  
work in progress....

# Conclusions

- Strongly interacting quantum critical systems are highly collective states without long-lived quasiparticles.
- Holography gives us examples of quantum critical states which behave like hydrodynamic fluids with a minimal viscosity.
- If a charged **hydrodynamic state with minimal viscosity** is weakly coupled to disorder/lattice, it will get a viscous contribution to its resistivity  $\rho_{DC}(T) \sim s(T)$ .
- This mechanism does not require holography. It may explain some of the strange transport properties of the strange metal phase of the high  $T_c$  cuprate superconductors.