How small hydrodynamics can go

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(Received 17 October 2020; accepted 16 February 2021; published 1 April 2021)

Numerous experimental and theoretical results in liquids and plasmas suggest the presence of a critical momentum at which the shear diffusion mode collides with a nonhydrodynamic relaxation mode, giving rise to propagating shear waves. This phenomenon, labeled "k-gap," could explain the surprising identification of a low-frequency elastic behavior in confined liquids. More recently, a formal study of the perturbative hydrodynamic expansion showed that critical points in complex space, such as the aforementioned k-gap, determine the radius of convergence of linear hydrodynamics—its regime of applicability. In this work, we combine the two new concepts, and we study the radius of convergence of linear hydrodynamics in "real liquids" by using several data from simulations and experiments. We generically show that the radius of convergence increases with temperature and it surprisingly decreases with the electromagnetic interactions coupling. More importantly, for all the systems considered, we find that such a radius is set by the Wigner–Seitz radius—the characteristic interatomic distance of the liquid, which provides a natural microscopic bound.

DOI: 10.1103/PhysRevD.103.086001

I. INTRODUCTION

" $\Pi \alpha \nu \tau \alpha \rho \epsilon i$ "—Everything flows. Hydrodynamics is an effective field theory (EFT) formulated as a perturbative expansion in spatial and time gradients. It governs the dynamics of conserved quantities which in Fourier space can be constructed as an infinite expansion in frequency ω and momentum k—from slow processes and large scales to fast dynamics and short lengths. It applies to the most disparate systems, from liquids [1] and solids [2] to flocks [3], crowds [4], and even financial markets [5]. As in every EFT, the microscopic physics is hidden in an infinite set of unknown coefficients since it is "irrelevant" (in the renormalization group sense) in the low energy regime of interest, where the EFT applies.

In this broad sense, we only assume the existence of local thermodynamic equilibrium and of small (or, rather, slow) fluctuations around it, whose dynamics is indeed described by hydrodynamics. Here, all the questions lie behind the words "small" and "slow" and their precise definitions and meanings.

In its linearized version, hydrodynamics is described by a finite set of *hydrodynamic modes*, which can be obtained from the knowledge of the conservation equations and the constitutive relations. These modes display dispersion relations which satisfy the requirement

$$\lim_{k \to 0} \omega_i(k) = 0. \tag{1}$$

Typical examples are diffusive modes $\omega = -iDk^2$ and propagating sound modes $\omega = \pm vk - i\Gamma k^2$. More broadly, every hydrodynamic mode obeys a dispersion relation,

$$\omega^{(i)}(k) = \sum_{n=1}^{\infty} \alpha_n^{(i)} k^n, \qquad (2)$$

where $\omega \in \mathbb{C}$ and $k \in \mathbb{R}$. These modes are practically obtained from an eigenvalues equation of the form

$$\prod_{j=1}^{N} (\omega - \omega^{(j)}(k)) \equiv F(\omega, k^2) = 0, \qquad (3)$$

which in general not only contains hydrodynamic modes but nonhydrodynamics¹ ones as well.

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¹Not satisfying Eq. (1).

By staring at Eq. (2), one could immediately ask if such a perturbative series is convergent and if yes which is its radius of convergence. This is tantamount to asking what is the radius of convergence of linearized hydrodynamics (in momentum space)—its regime of applicability.² Reference [8] and later on, in a more complete form, Refs. [9,10] suggested that in order to answer such question, one has to formally extend the function $F(\omega, k^2)$ in complex momentum space and treat it as a complex algebraic curve. In this language, series like Eq. (2) are known as Puiseux series, and their radius of convergence is fundamentally connected to the so-called *critical points* { ω_c, k_c }—points at which

$$F(\omega_c, k_c^2) = 0, \qquad \frac{\partial F(\omega_c, k_c^2)}{\partial \omega} = 0,$$
 (4)

for both ω_c , $k_c^2 \in \mathbb{C}$. These critical points signal "irregularities" in the algebraic curve, and they are related to quasinormal modes (QNMs)³ crossing. The radius of convergence \mathcal{R} of the hydrodynamic expansion is then set by the distance to the first of these critical points:

$$\mathcal{R} \equiv |k_c|. \tag{5}$$

Moreover, the radius of convergence is intimately connected to the existence of nonhydrodynamic excitations and their interactions with the hydrodynamics modes.

This mathematical machinery is extremely elegant but also rather abstract and hard to digest. So far, it has been applied only to few holographic models [12,13] whose relevance for more realistic situations is at least disputable. In this work, we address the question of the regime of applicability of hydrodynamics in "real" liquids, and in particularly we ask the following question: given a specific liquid, until which length scale are we allowed to trust the hydrodynamic approximation? This question can be rephrased as follows: "Which is the physical (and not mathematical) scale setting the breakdown of the hydrodynamic expansion?" A naive answer would be that hydrodynamics is a good description of a liquid as far as the full system can be seen as a continuum and not as a set of molecules/particles interacting with each others. Technically, the failure of hydrodynamics is related to the intrusion of nonhydrodynamic modes which cannot be neglected anymore.

Combining the mathematical methods of Refs. [9,10] with data from experiments and molecular dynamics (MD) simulations, we will indeed show that the limiting length

scale for the hydrodynamic framework is given by an $\mathcal{O}(1)$ fraction of the intermolecular distance, given formally by the size of Wigner-Seitz cell *a*. We will also discuss how the regime of applicability depends on the temperature *T* and on the interactions strength. As we will discuss in detail, the reinterpretation of the available data will lead us to confirm some intuitive physical arguments but also to new and unexpected findings such as the fact that hydrodynamics *does not* work better at strong electromagnetic coupling (as always advertised).

II. CONVERGENCE OF LINEAR HYDRODYNAMICS IN LIQUIDS

For centuries, the presence of low-frequency propagating shear waves (i.e., transverse phonons) has been considered the fundamental criterion to distinguish solids from liquids. Nevertheless, in the last decade, this definition has been challenged by several experiments observing the presence of shear waves in confined liquids at small frequency [14] and corresponding solidlike elastic effects [15,16] (see also Ref. [17]). This phenomenon indicates that the difference between solids and liquids is only quantitative and it is measured by the relaxation time τ —the average time for molecules rearrangement. Within Maxwell theory, such a timescale is simply $\tau \equiv \eta/G$ with η the shear viscosity and G the shear elastic modulus. It follows that in a solid $\tau = \infty$ —molecules do not rearrange but just oscillate around their equilibrium positions. From a theoretical point of view, this program goes under the name of Maxwell interpolation, and its main result is that the dynamics of shear waves in a fluid is governed by the simple equation

$$\omega^2 + i\omega/\tau - v^2 k^2 = 0,$$
 (6)

which is known for historical reason as the *telegraph* equation. Here, v is the asymptotic speed of the shear waves, dictated by the elastic modulus. Solving Eq. (6), one finds a couple of modes,

$$\omega = -\frac{i}{2\tau} \pm \sqrt{v^2 k^2 - \frac{1}{4\tau^2}} \tag{7}$$

and the existence of a critical momentum k_g , labeled as k_{gap} , at which transverse waves start to propagate (Re $\omega \neq 0$). This mechanism appears in several physical contexts; we refer the reader to Ref. [18] for a complete review of them.

The appearance of propagating waves for $k > k_g$ is the consequence of the collision between the shear diffusion mode and a nonhydrodynamic excitation, which happens exactly at $k = k_g$ (see Fig. 1). Given Eq. (6), it is straightforward to apply the methods of Refs. [9,10] and determine the critical point of the corresponding algebraic curve using Eq. (4). One immediately obtains that $\omega_c = -i/2\tau$ and more importantly that

²In this work, we will not consider the question of convergence in real space and for the full nonlinear dynamics. For that, see for example Ref. [6] and specially Ref. [7].

³By these, we refer simply to all modes with dispersion relation $\omega(k)$ with $k \in \mathbb{R}$. In other words, QNMs are all the excitations of a specific system [11].



FIG. 1. The lowest collective modes in the transverse sector of a liquid, following from the k-gap equation (6). The real part of the dispersion relation is displayed with a dashed line, and the imaginary is displayed with a solid line. We set $v = \tau = 1$ for which the cutoff momentum is $k_q = 0.5$.

$$\mathcal{R} \equiv |k_c| = \frac{1}{2v\tau} = k_g. \tag{8}$$

This result is confirmed by explicitly computing the set of algebraic curves in the complex plane for complex momentum (see Fig. 2). Notice that this already implies



$|\omega_c|\tau = \frac{1}{2},\tag{9}$

momentum $L \equiv k_g^{-1}$. Before proceeding, it is important to discuss the assump-

tions behind the telegraph equation (6) and its validity. Clearly, this quadratic equation is a good description of the low energy dynamics only when the system displays a separation of scales between the two lower excitations considered in Eq. (6) and the rest of the modes [i.e., higher order corrections in (6)]. This depends *a priori* on the microscopic details of the system, and it can be for instance achieved whenever the relaxation time τ is parametrically long (e.g., in presence of a weakly broken symmetry) and the quasihydrodynamics description of Ref. [19] is valid. Two more comments are in order:

meaning that linearized hydrodynamics breaks down for a

timescale shorter than twice the intrinsic relaxation time. In summary, in realistic liquids, in which the dynamics of

shear waves is well described by the k-gap equation (6), hydrodynamics applies from large distances until a "microscopic" length scale given by the inverse of the k-gap

(i) In our analysis, we do not rely at any time on the specific relation given by the telegraph equation $k_g = 1/(2v\tau)$. Higher momentum corrections obviously modify such expression; nonetheless, the radius of convergence would still be given by the collision point and well approximated by the value at which the real part of the dispersion relation becomes nonzero.

(ii) The numerical dispersion relations from which the data presented are extracted are very well fitted by the square-root k-gap expression coming from Eq. (6), at least in the vicinity of the critical momentum of interest. This is an *a posteriori* proof that the potentially dangerous higher order corrections are indeed negligible and that higher modes do not participate in this dynamics. In summary, within the regimes discussed, our analysis is expected to be robust, independent of the validity of the simplex Maxwell model and affected by higher order corrections only in a minor way.

III. DATA FROM EXPERIMENTS AND SIMULATIONS

The validity of the k-gap equation (6) to describe the shear dynamics in liquids has been extensively corroborated by several MD simulations and experiments (EXP) in two-dimensional (2D) and three-dimensional (3D) liquids and plasmas [20–29]. The dispersion relation of the transverse waves is usually extracted from the transverse current correlation function $\langle J(-k, t)J(k, 0) \rangle$ and within the numerical/experimental errors is well fitted by

$$\operatorname{Re}\omega = \sqrt{v^2 k^2 - \frac{1}{4\tau^2}} \quad \text{for } k > k_g.$$
(10)

It is therefore obvious how to extract the critical momentum from the data.

FIG. 2. The complex curves corresponding to Eq. (6) for complex momentum $k = |k|e^{i\theta}$ with $\theta \in [0, 2\pi]$ and varying the modulus |k|. The magenta curve corresponds to the critical momentum $|k_c| = 0.5$. We set $v = \tau = 1$.

Unfortunately, the critical point determining the radius of convergence in the longitudinal sector is not located at real values of the momentum k. For this reason, we are not in the position to complete this analysis including the longitudinal sector. To the best of our knowledge, it is at the moment impossible to obtain data for complex momentum (and complex frequency). Moreover, a complete understanding of the longitudinal sound dynamics in liquids (e.g., the "positive sound dispersion" phenomenon [30]), analogous to the k-gap phenomenon for shear waves, is still elusive. A priori, one cannot determine whether the constraint on convergence coming from the longitudinal sector would be more stringent than the one derived in this work. Nevertheless, from a simple physical argument, one would expect both scales to be controlled by the only microscopic length scale in liquids-the intermolecular distance.

A fundamental parameter, both in the simulations and in the experiments, is given by the intermolecular distance *a*, which is extracted from the position of the first maximum in the pair distribution function. Technically, this scale corresponds to the Wigner-Seitz radius $a = \left(\frac{2(D-1)}{D}n\right)^{-1/D}$, where *n* is the number density and *D* is the number of spatial dimensions. In Table I, we list the value of the product $k_c a$ for a large number of 2D and 3D liquids and plasma, and for all the systems considered, we do find that

$$k_c a \approx \mathcal{O}(1). \tag{11}$$

This result robustly indicates that the radius of convergence of hydrodynamics is set by the intermolecular distance.⁴ Moreover, it is in agreement with the naive idea that hydrodynamics applies up to the scale at which you can resolve that the continuum of the liquid is actually formed by a collection of particles. It is important to notice that the relation (11) is highly nontrivial since the Wigner-Seitz radius can vary from Å (10^{-10} m) for most of the fluids to millimeter scale in dusty plasmas (see for example Ref. [23]).

Interestingly, the critical momentum for quark gluon plasma was estimated in Ref. [31] around $k_c^{-1} \approx 0.15$ fm. Assuming an interparton distance of the order of $a \approx$ 0.5 fm [32], we again obtain a value $k_c a = \mathcal{O}(1)$. This fact is reminiscent of the universality found in the values of the kinematic viscosity in Ref. [33]. As a comment, it would be interesting to understand which quantity plays the role of the microscopic intermolecular scale *a* in a liquid which appears to be strongly coupled at all energy scales (if it exists).

TABLE I. A summary of the available data for the momentum cutoff of shear waves k_c in liquids and plasma. The interatomic distance is defined as a.

Liquid	k _c a
2D Yukawa (MD) [20]	0.25
Dusty plasma (MD) [21,22]	0.3-1.2
2D Yukawa (EXP) [23]	0.16-0.31
Liquid Fe (EXP) [24,25]	0.3
Liquid Cu (EXP) [24,25]	0.4
Liquid Zn (EXP) [24,25]	0.3
3D Lennard-Jones fluid (MD) [26]	0.2-0.7
Liquid Fe (MD) [26]	0.2-0.7
IPL8-IPL12 fluid (MD) [26]	0.2-0.7
Liquid Hg (MD) [26]	0.15-0.55
Supercritical Ar (MD) [27]	0.05-0.8
Subcritical liquid Ar (MD) [27]	0.2-0.7
Supercritical CO ₂ [27]	0.1-0.5
Liquid Ga (EXP, MD) [28]	0.25-0.6
2D Coulomb classical fluids (MD) [29]	0.3-2
Quark gluon plasma [31]	3.3

Given the abundance of available data, we can go one step further and investigate the temperature dependence of the hydrodynamic convergence radius. In Fig. 3, we have plotted the data for nine different liquids in a large range of temperatures. In all the cases, the regime of applicability increases with temperature, and it is consistent with the idea that $k_g \sim 1/v\tau$ and τ decreases monotonically with temperature (in standard liquids following the well-known Arrhenius law). It also connects with the idea that at T=0(or equivalently $\tau = \infty$) hydrodynamics is not applicable.⁵

Finally, using the data for Coulomb fluids and plasmas, we can also investigate the radius of convergence in terms of the effective coupling parameter,

$$\Gamma \equiv \frac{Q^2}{4\pi\epsilon_0 a k_B T},\tag{12}$$

which determines the strength of the Coulomb interactions in the plasma. Here, Q is the charge, and ϵ_0 is the dielectric constant. The data from simulations universally show (see Fig. 4) a powerlike decrease of the critical momentum in function of Γ , which is further confirmed by experimental data [23]. The dusty plasma considered is a suspension of highly charged microsize particles which repel each others via a nearly Coulomb potential,

$$U(r) = \frac{Q}{4\pi\epsilon_0 r} e^{-r/\lambda_D},\tag{13}$$

in which the exponential correction takes into account the screening effects via the screening length λ_D . The results therefore demonstrate that stronger Coulomb repulsion

⁴In principle, the critical scale k_c could be pushed all the way down to zero by achieving a very large relaxation time τ (i.e., in glasses or solids). Nevertheless, in that situation, the hydrodynamic window $\omega \tau \ll 1$ would shrink completely, and therefore hydrodynamics would not be applicable at all. We thank Kostya Trachenko for pointing this out.

⁵This follows simply by the violation of the requirements $\omega/T \ll 1$ or $\omega \tau \ll 1$.



FIG. 3. The dimensionless combination $k_c a$ with $k_c = k_g$ determining the radius of convergence and *a* the intermolecular characteristic distance. Temperature dependence for IPL8 fluid (black) [26], IPL12 fluid (gray) [26], liquid supercritical argon sAr (yellow) [27], supercritical CO₂ (green) [27], supercritical argon sAR (brown) [27], liquid Ga (orange) [28], liquid Hg (red) [26], liquid Fe $\rho = 8$ gm/cm³ (purple) [26], and liquid Fe $\rho = 10$ g/cm³ (blue) [26].



FIG. 4. The dimensionless combination $k_c a$ with $k_c = k_g$ determining the radius of convergence and *a* the intermolecular characteristic distance. The empty squares indicate data from MD simulations taken from Refs. [21,25,29]. The filled squares are experimental data from Ref. [23]. See also Ref. [34] for a similar collection of data.

decreases the regime of applicability of hydrodynamics. This observation seems to contradict the common wisdom that hydrodynamics works better for strongly coupled systems in which the mean free path becomes smaller (see for example Fig. 4 in Ref. [31] and Appendices A and B). Nevertheless, it is important to notice that the coupling therein refers to the nuclear interactions—the strength of the bound states [corresponding to the SU(N) gauge group] —and not to the intermolecular Coulomb force.⁶ Therefore, no evident discordance is present.

IV. CONCLUSIONS

In this work, we have combined the theoretical methods proposed by Refs. [9,10] with a large set of data from molecular dynamics simulations and experiments in 2D and 3D liquids and plasmas to determine the regime of applicability of hydrodynamics in realistic systems—the hydrodynamic convergence radius. Our main result is that the convergence radius is always set in terms of the intermolecular distance, namely the length scale at which one can resolve the independent set of particles within the fluid continuum.

Moreover, the data indicate that the regime of validity of hydrodynamics increases with the temperature T and decreases with the effective electromagnetic coupling parameter Γ . This trend is totally unexpected and against the usual (nevertheless never demonstrated) slogan that "hydrodynamics works better at strong coupling." This point definitely deserves further investigation.

Interestingly, the critical momentum for plasmon modes⁷ in Dirac fluids has been recently computed in Ref. [35], and it reads

$$k_c = \frac{\tau_V}{4v\tau_{c,1}^2},\tag{14}$$

where $\tau_{c,1}$ is the relaxation time, v is the electrons group velocity, and τ_V is the timescale characterizing electrostatic interaction. We are planning to perform a more detailed comparison between Eq. (14) and our results in the near future.

Taking into account the fundamental role of nonhydrodynamic modes, it would be extremely helpful to build experimental setups able to pinpoint them on the lines of Ref. [36]. Moreover, it would be interesting to understand which are the experimental and physical consequences of the nonconvergence of the hydrodynamic series in realistic systems.⁸

Finally, the k-gap dynamics (6) universally appears in the context of diffusive Goldstone bosons [37,38] in dissipative systems (e.g., in quasicrystals [39,40]). It would be interesting to connect such mechanism with the results of this paper.

In conclusion, given the extremely wide usage of hydrodynamic methods, it is mandatory to understand until which length scale those can be trusted. In this work, we provided a direct and pragmatic answer to this question in a large set of realistic liquids and plasmas. We hope that our results will boost the efforts to deeply understand hydrodynamics and its regime of applicability and to connect more closely theory with experiments.

⁶We thanks Saso Grozdanov for suggesting this point.

⁷Plasmons modes are not QNMs, but they are solution of the equation $\epsilon(\omega, k)$ where ϵ is the dielectric constant. The latter can be taken as our abstract function $F(\omega, k^2)$ in (4).

We thank Egor Kiselev for suggesting this point.

ACKNOWLEDGMENTS

The author would like to thank A. Zaccone, S. Yurchenko, L. Noirez, N. Poovuttikul, N. Abbasi, S. Grieninger, M. Heller, J. Schmalian, E. Kiselev, K. Landsteiner, and specially Grozdanov, S. P. Romantschke, and K. Trachenko for fruitful discussions and insightful comments on the topics covered in this manuscript. The author acknowledges the support of the Shanghai Municipal Science and Technology Major Project (Grant No. 2019SHZDZX01) and of the Spanish MINECO "Centro de Excelencia Severo Ochoa" Programme under Grant No. SEV-2012-0249.

APPENDIX A: HOLOGRAPHIC EXAMPLES

A common playground where the k-gap equation (6) appears is in the context of holographic models with higher-from global symmetries [19,41–43]. These models are used to described both magneto-hydrodynamics in charged plasmas and the elasto-dynamics in viscoelastic media. In the first scenario, the emergent propagating degree of freedom at $k = k_g$ is the photon. In the neutral plasma approximation, the corresponding critical momentum is given in terms of [44]

$$k_c \approx \frac{\pi T}{2\log(M/\pi T)},\tag{A1}$$

where T is the temperature and M is an UV cutoff of the theory.

In the second scenario, the emergent mode is a propagating phonon and the critical momentum reads [45]





FIG. 5. The critical momentum in the higher-form global symmetries holographic models. The formulas are taken from [19,42] for the photon propagator and from [43] for the viscoelastic model.



FIG. 6. The critical momentum in function of the electromagnetic coupling λ in the holographic model of [47].

where the same notations are used. It is simple to see that in both cases the critical momentum, determining the radius of convergence of hydrodynamics, grows monotonically with temperature independently of the value of the UV cutoff M as shown in Fig. 5.

Another situation where the k-gap equation (6) is at work is in models with dynamical Coulomb interactions and emergent plasmonic modes [46–48]. In those models, it is possible to obtain numerically the dispersion relations in function of the electromagnetic coupling λ . The numerical results are shown in Fig. 6 and they clearly indicate a power-law fall-off of the critical momentum in function of λ . This is in agreement with the data obtained from 2D plasmas and presented in the main text (see Fig. 4) under identifying λ with Γ .

Finally, a well-known place where the k-gap appears is in the Israel-Stewart theory for linearised relativistic hydrodynamics [49]. In this model, the critical momentum is given by

$$k_c = \frac{1}{\sqrt{4D_\tau}} = \sqrt{\frac{\epsilon + p}{4\eta\tau_\pi}}.$$
 (A3)

The gravity dual of the Israel-Stewart theory is provided by a Gauss-Bonnet model [19], where the k-gap momentum reads:

$$k_c = -\frac{\sqrt{2}\pi T \sqrt{-\frac{\lambda_{\rm GB}}{\gamma(\gamma+2)+2\log(\frac{2}{\gamma+1})-3}}}{\lambda_{\rm GB}},\qquad({\rm A4})$$

with $\gamma \equiv \sqrt{1 - 4\lambda_{GB}}$ and $\lambda_{GB} \in [-\infty, 1/4]$. From Eq. (A4) two facts are evident. (I) The critical momentum grows linearly with temperature in agreement with the data shown in Fig. 3. (II) The critical momentum decreases by increasing the value of the Gauss-Bonnet coupling on the negative axes. Given that such direction corresponds to decrease the coupling of the dual field theory from infinity, the results shown in Fig. 7 suggest that



FIG. 7. The critical momentum $k_c = 1/2v\tau$ in function of the Gauss Bonnet coupling. Formulas taken from [19].

hydrodynamics works better at strong coupling. This outcome is in agreement with [31]. Notice that the disagreement with the behavior with respect to Γ is certainly puzzling but it does not constitute a discordance since λ refers to the SU(N) coupling while Γ to the inter-molecular Coulomb potential.

APPENDIX B: THE TELEGRAPH

As explained in the main text, Eq. (6) is also known as the *telegrapher equation* and it was first written by Oliver Heaviside in the attempt of studying the dynamics of transmission lines. In particular, such equation comes from the analysis of a specific circuit displayed in Fig. 8, where R, L, G, C are respectively a resistance, an inductance, a conductance and a capacitance. In such a configuration, the equation for the voltage v across the piece of wire at position x at time t is given by:

$$LC\frac{\partial^2 v}{\partial t^2} + (LC + RG)\frac{\partial v}{\partial t} + RGv = \frac{\partial^2 v}{\partial x^2}, \quad (B1)$$



FIG. 8. Schematic of the electric circuit giving rise to the telegraph equation Eq. (B1).

which is a generalized form of Eq. (6) with the presence of an extra mass term (vanishing for either R = 0 or G = 0). In any case, the solution enjoys very similar properties and in particular the presence of a critical momentum defined as

$$k_c = \frac{|GL - CR|}{2\sqrt{CL}}.$$
 (B2)

Interestingly, using some data of telegraph wires for transmissions at low frequency reported in [50], one obtains:

$$k_c \approx 0.79 \text{ Km}^{-1}, \tag{B3}$$

which means that the radius of convergence of hydrodynamics in a telegraph wire is about $L \approx 1.26$ Km! Moreover, one could easily extract the relaxation time:

$$\tau = \frac{CL}{GL + CR} \approx 10^{-6} \text{ s} \tag{B4}$$

which determines the timescale at which hydro fails. Notice that obviously:

$$\frac{1}{k_c \tau} \approx 3 \times 10^8 \text{ m/s} \equiv c, \tag{B5}$$

i.e., the speed of light—the speed of propagation of electromagnetic waves inside the transmission wire.

- L. Landau and E. Lifshitz, *Fluid Mechanics*, Course of Theoretical Physics S Vol. 6 (Elsevier Science, New York, 2013).
- [2] P. C. Martin, O. Parodi, and P. S. Pershan, Phys. Rev. A 6, 2401 (1972).
- [3] J. Toner, Y. Tu, and S. Ramaswamy, Ann. Phys. (Amsterdam) 318, 170 (2005).
- [4] N. Bain and D. Bartolo, Science 363, 46 (2019).
- [5] Y. Yura, H. Takayasu, D. Sornette, and M. Takayasu, Phys. Rev. Lett. **112**, 098703 (2014).
- [6] M. P. Heller, A. Serantes, M. Spaliński, V. Svensson, and B. Withers, arXiv:2007.05524.
- [7] P. Romatschke and U. Romatschke, *Relativistic Fluid Dynamics In and Out of Equilibrium*, Cambridge Monographs on Mathematical Physics (Cambridge University Press, Cambridge, England, 2019).
- [8] B. Withers, J. High Energy Phys. 06 (2018) 059.
- [9] S. C. v. Grozdanov, P. K. Kovtun, A. O. Starinets, and P. Tadić, Phys. Rev. Lett. 122, 251601 (2019).
- [10] S. Grozdanov, P. K. Kovtun, A. O. Starinets, and P. Tadić, J. High Energy Phys. 11 (2019) 97.
- [11] P. K. Kovtun and A. O. Starinets, Phys. Rev. D 72, 086009 (2005).
- [12] N. Abbasi and S. Tahery, J. High Energy Phys. 10 (2020) 076.

- [13] A. Jansen and C. Pantelidou, J. High Energy Phys. 10 (2020) 121.
- [14] L. Noirez and P. Baroni, J. Phys. Condens. Matter 24, 372101 (2012).
- [15] E. Kume, P. Baroni, and L. Noirez, Sci. Rep. 10, 13340 (2020).
- [16] E. Kume, A. Zaccone, and L. Noirez, arXiv:2009.09788.
- [17] M. Grimsditch, R. Bhadra, and L. M. Torell, Phys. Rev. Lett. 62, 2616 (1989).
- [18] M. Baggioli, M. Vasin, V. Brazhkin, and K. Trachenko, Phys. Rep. 865, 1 (2020).
- [19] S. C. V. Grozdanov, A. Lucas, and N. Poovuttikul, Phys. Rev. D 99, 086012 (2019).
- [20] G. J. Kalman, P. Hartmann, Z. Donkó, and M. Rosenberg, Phys. Rev. Lett. 92, 065001 (2004).
- [21] M.S. Murillo, Phys. Rev. Lett. 85, 2514 (2000).
- [22] H. Ohta and S. Hamaguchi, Phys. Rev. Lett. 84, 6026 (2000).
- [23] V. Nosenko, J. Goree, and A. Piel, Phys. Rev. Lett. 97, 115001 (2006).
- [24] S. Hosokawa, M. Inui, Y. Kajihara, S. Tsutsui, and A. Q. R. Baron, J. Phys. Condens. Matter 27, 194104 (2015).
- [25] S. A. Khrapak, A. G. Khrapak, N. P. Kryuchkov, and S. O. Yurchenko, J. Chem. Phys. **150**, 104503 (2019).
- [26] N. P. Kryuchkov, L. A. Mistryukova, A. V. Sapelkin, V. V. Brazhkin, and S. O. Yurchenko, Phys. Rev. Lett. 125, 125501 (2020).
- [27] C. Yang, M. T. Dove, V. V. Brazhkin, and K. Trachenko, Phys. Rev. Lett. 118, 215502 (2017).
- [28] R. M. Khusnutdinoff, C. Cockrell, O. A. Dicks, A. C. S. Jensen, M. D. Le, L. Wang, M. T. Dove, A. V. Mokshin, V. V. Brazhkin, and K. Trachenko, Phys. Rev. B 101, 214312 (2020).
- [29] S. A. Khrapak, N. P. Kryuchkov, L. A. Mistryukova, A. G. Khrapak, and S. O. Yurchenko, J. Chem. Phys. 149, 134114 (2018).
- [30] Y. H. Jeong, S. R. Nagel, and S. Bhattacharya, Phys. Rev. A 34, 602 (1986).
- [31] P. Romatschke, Eur. Phys. J. C 77, 21 (2017).

- [32] E. Shuryak, Rev. Mod. Phys. 89, 035001 (2017).
- [33] K. Trachenko, V. Brazhkin, and M. Baggioli, arXiv:2003 .13506.
- [34] J. Goree, Z. Donkó, and P. Hartmann, Phys. Rev. E 85, 066401 (2012).
- [35] E. I. Kiselev and J. Schmalian, Phys. Rev. B 102, 245434 (2020).
- [36] J. Brewer and P. Romatschke, Phys. Rev. Lett. 115, 190404 (2015).
- [37] Y. Minami and Y. Hidaka, Phys. Rev. E **97**, 012130 (2018).
- [38] Y. Hidaka and Y. Minami, Prog. Theor. Exp. Phys. 033A01 (2020).
- [39] R. Currat, E. Kats, and I. Luk'yanchuk, Eur. Phys. J. B 26, 339 (2002).
- [40] M. Baggioli and M. Landry, SciPost Phys. 9, 062 (2020).
- [41] S. Grozdanov, D. M. Hofman, and N. Iqbal, Phys. Rev. D 95, 096003 (2017).
- [42] D. M. Hofman and N. Iqbal, SciPost Phys. 4, 005 (2018).
- [43] S. C. v. Grozdanov and N. Poovuttikul, Phys. Rev. D 97, 106005 (2018).
- [44] S. Grozdanov and N. Poovuttikul, J. High Energy Phys. 04 (2019) 141.
- [45] S. Grozdanov, N. Kaplis, and A.O. Starinets, J. High Energy Phys. 07 (2016) 151.
- [46] U. Gran, M. Tornsö, and T. Zingg, J. High Energy Phys. 2018 (2018) 176.
- [47] M. Baggioli, U. Gran, A. J. Alba, M. Tornsö, and T. Zingg, J. High Energy Phys. 2019 (2019) 13.
- [48] M. Baggioli, U. Gran, and M. Tornsö, J. High Energy Phys. 04 (2020) 106.
- [49] W. Israel and J. Stewart, Ann. Phys. (N.Y.) 118, 341 (1979).
- [50] W. Chen, Home Networking Basis: Transmission Environments and Wired/wireless Protocols, Communications Engineering (Prentice Hall PTR, NJ, 2004), https://www .amazon.com/Home-Networking-Basis-Transmission-Environments/dp/0130165115.