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Proposal for universality in the viscosity of metallic liquids

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The range of magnitude of the liquid viscosity, η , as a function of temperature is one of the most impressive of any physical property, changing by approximately 17 orders of magnitude from its extrapolated value at infinite temperature (η_0) to that at the glass transition temperature, T_g . We present experimental measurements of containerlessly processed metallic liquids that suggest that $\log(\eta/\eta_0)$ as a function of T_A/T is a potentially universal scaled curve. In stark contrast to previous approaches, the scaling requires only two fitting parameters, which are on average predictable. The temperature T_A corresponds to the onset of cooperative motion and is strongly correlated with T_g , suggesting that the processes underlying the glass transition first appear in the high temperature liquid.

The nature of the dynamical processes in liquids and how liquids transform to glasses are major outstanding questions in condensed matter science. The shear viscosity is a particularly temperature-sensitive property for glass-forming liquids, changing by about 17 orders of magnitude upon cooling from high temperatures to the glass transition temperature, T_g . The way in which the viscosity, or related relaxation times, change with temperature scaled to T_g is the basis for the widely-used *fragility* classification scheme introduced by Angell¹. For very *strong* liquids the viscosity shows an Arrhenius behavior, with a well-defined activation energy over a wide temperature range that extends from above the melting temperature down to T_g . The viscosities of fragile liquids are characterized by activation energies that are small at high temperature and increase rapidly upon approaching T_g . The strongest glass-formers are network oxides, while molecular liquids such as o-terphenyl, decalin and isoquinoline, are among the most fragile. Upon close examination, some non-Arrhenius behavior is observed near the glass transition, even in strong liquids, but this becomes more dramatic as the fragility of the liquid increases. Thermodynamic² and direct structural signatures³ of fragility support a connection between structure and dynamics in liquids, which has been long assumed. The concept of fragility appears to provide a coherent scheme for classifying all liquids and linking to glass formability in some cases. However, in spite of the significant variation in fragility among various liquid metals, the viscosity experimental data for those presented here are remarkably more universal. They suggest the existence of a high-temperature universal scaling temperature, T_A , which has been predicted from molecular dynamics (MD) simulations of metallic liquids⁴ and theoretical studies of non-metallic glass-forming liquids^{5–7}. Scaling the temperature, T , by T_A (a temperature that corresponds to the onset of dynamical cooperativity in the liquid), and scaling the viscosity by η_0 (the extrapolated high temperature limit on the viscosity of the liquid phase) yields an apparently universal curve that fits the viscosities of all liquid metals studied, from above the melting temperature to the glass transition temperature.

Results

Measurements of the Liquid Viscosity and a Universal Curve. The viscosities of a variety of metallic liquids were measured at high temperatures in a high-vacuum containerless environment, using electrostatic levitation (see Supplementary Information). These are shown as a function of inverse temperature in Fig. 1A. The liquids studied include liquids that easily form metallic glasses, such as $Zr_{57}Cu$

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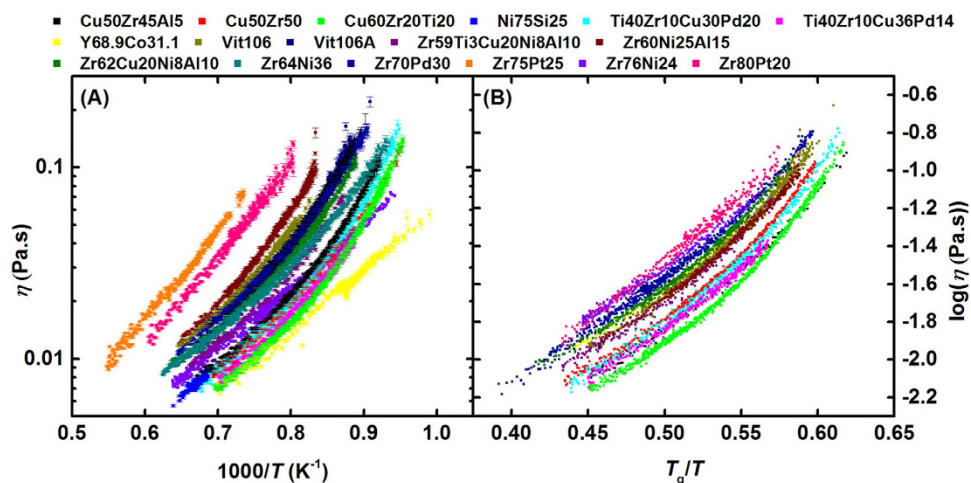


Figure 1. (A) Measured viscosity, η , as a function of temperature for metallic liquids. Error bars are one s.d. (B) An Angell plot of the $\log(\eta)$ as a function of T_g/T .

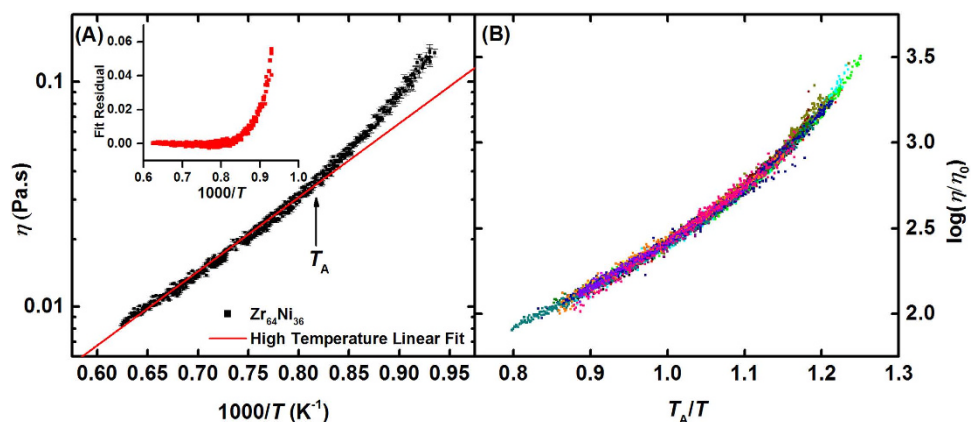


Figure 2. (A) Typical example of the behavior of $\log(\eta)$ at high temperature, showing a departure from Arrhenius behavior on cooling below T_A . Error bars are one s.d. (B) Scaled universal curve for all measured viscosity data.

$_{15.4}\text{Ni}_{12.6}\text{Al}_{10}\text{Nb}_5$ (Vit 106) and $\text{Zr}_{58.5}\text{Cu}_{15.6}\text{Ni}_{12.8}\text{Al}_{10.3}\text{Nb}_{2.8}$ (Vit 106A), and more marginal glass-forming fragile liquids, such as Ti-Zr-Cu-Pd and Cu-Zr, where faster cooling rates are required for glass formation. Also included are liquids for which glass formation has not been observed, such as Ni-Si. Figure 1B shows an Angell plot for these data, presenting $\log(\eta)$ as a function of T_g/T . Scaling the temperature by T_g reduces the scatter of the data from Fig. 1A, but significant variations among the data remain.

As illustrated in Fig. 2A for a $\text{Zr}_{64}\text{Ni}_{36}$ liquid, the viscosities of all of the data in Fig. 1 have Arrhenius temperature dependences at a sufficiently high temperature. This agrees with the results from previous studies^{5,8,9} and recent MD simulations for several different types of metallic liquids⁴. The temperature at which the measured viscosity departs from Arrhenius behavior is labeled in Fig. 2A as T_A . While the departure is gradual and difficult to determine directly, it becomes clearer in the residuals from a linear fit (insert to figure). The physical meaning of T_A is intriguing; from MD simulations⁴ it corresponds to the temperature at which flow first becomes cooperative.

The MD simulations revealed a universal curve for the ratio of the Maxwell relaxation time for viscosity and the time required to change the local coordination number in a cluster by one unit, by scaling the temperature with the temperature corresponding to the onset of dynamical cooperativity (defined there also as T_A). Those results suggested that the measured viscosity data could be scaled by T_A . More recent simulation results reveal concurrent cooperative structural changes beginning at T_A ¹⁰. As is verified in Fig. 2B, by adopting two material-dependent parameters for scaling, T_A and η_0 , all of the data can be collapsed into a universal curve that describes the temperature dependent viscosity of the liquids studied. To construct this curve, the scaling temperature, T_A , was determined for each liquid as illustrated in Fig. 2A, defined as the temperature below which the viscosity became non-Arrhenius. The value of η_0 for each

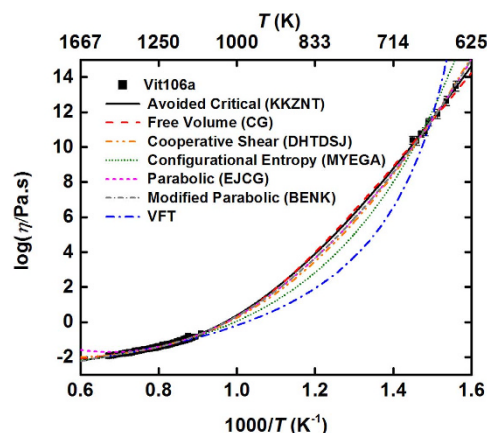


Figure 3. Comparison of the fits to the Vit 106a data for the expressions listed in Table 1. The fit parameters can be found in Supplementary Table 1. Error bars are one s.d.

Vogel-Fulcher-Tammann (VFT)	$\log \eta = \log \eta_0 + \frac{D^* T_0}{T - T_0}$
Configurational Entropy (MYEGA)	$\log \eta = \log \eta_0 + \frac{K}{T} \exp\left(\frac{C}{T}\right)$
Free Volume (CG)	$\log \eta = \log \eta_0 + 2B/(T - T_0 + \sqrt{(T - T_0)^2 + CT})$
Avoided Critical (KKZNT)	$\log \eta = \log \eta_0 + \frac{1}{T}(E_\infty + T^* B[(T^* - T)/T^*]^z \Theta(T^* - T))$
Cooperative Shear (DHTDSJ)	$\log \eta = \log \eta_0 + \frac{W_0}{KT} \exp\left(-\frac{T}{T_W}\right)$
Parabolic (EJCG)	$\log \eta = \log \eta_0 + J^2 \left(\frac{1}{T} - \frac{1}{T_0}\right)^2, T < T_{FitMax} \leq T_0$
Modified Parabolic (BENK)	$\log \eta = \log \eta_0 + \frac{E}{KT} + J^2 \left(\frac{1}{T} - \frac{1}{\tilde{T}}\right)^2 \Theta(\tilde{T} - T)$

Table 1. Tested Fitting Functions for Viscosity. $\Theta(x)$ is the Heaviside function (i.e., $\Theta(x > 0) = 1$ and $\Theta(x < 0) = 0$).

liquid was then adjusted to collapse the data along the vertical axis. For all liquids studied, on average $\eta_0 \approx nh^{11,12}$, where n is the particle density and h is Planck's constant. Exact values and comparisons to nh can be found in Supplementary Table 2.

Functional Form of the Universal Curve. While it is shown that our measured data lie on a universal curve when properly scaled, the temperature range is small. An immediate consequence of the observed data collapse is that, at intermediate temperatures, $\eta = \eta_0 F(T/T_A)$. Here, $F(z)$ is assumed to be a universal function common to all of the metallic liquids studied (i.e., one exhibiting *no adjustable parameters*). To examine whether the scaling holds over a larger temperature range, measured viscosity data near T_g for the strong metallic glass-forming liquid Vit 106a¹³ were combined with the high temperature data reported here for the same liquid. However, since the scaling in Fig. 2B is empirical, extending it to lower temperatures requires knowledge of the functional form of the universal curve. Many earlier proposed expressions for the viscosity exist, but it has not been previously possible to extensively test them in metallic liquids over the wide temperature range that is possible here, extending from above the melting temperature to the glass transition temperature. The combined Vit 106a data, then, constitute a benchmark for testing these expressions and for identifying the one (if any) that describes the universal behavior. The fits to the Vit 106a data for some of the better-known expressions are shown in Fig. 3. These are (i) the commonly used Vogel-Fulcher-Tammann (VFT) equation¹⁴, (ii) the recently proposed Mauro-Yue-Elliston-Gupta-Allan (MYEGA) equation¹⁵, (iii) a relation derived within the Cohen-Grest free volume model (CG)¹⁶, (iv) the avoided critical point theory (KKZNT)⁵⁻⁷, (v) a cooperative shear model (DHTDSJ)¹⁷, and (vi) a parabolic kinetically constrained model (EJCG)^{18,19}. Since the EJCG expression is only valid up to its onset temperature T_0 , leaving higher temperature behavior undefined, a new variant (BENK) is introduced here, for which the constant Arrhenius type barrier is augmented to give to an expression that emulates that of the avoided critical point theory when T is replaced by $1/T$. Its crossover temperature \tilde{T} is similar to T^* in the KKZNT expression. These equations are defined in Table 1 and the corresponding optimal parameters are listed in Supplementary Table 1. Other approaches²⁰⁻²² that are associated with scaling temperatures typically near the mode-coupling temperature, below our experimentally accessible temperature region, could not be examined.

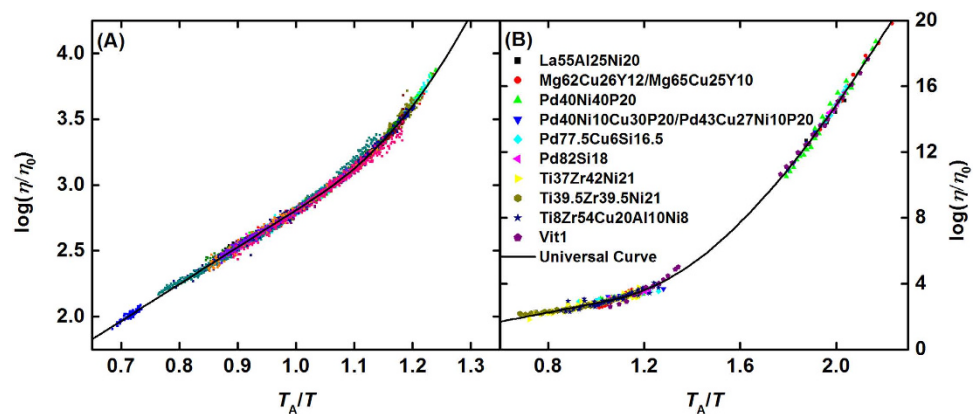


Figure 4. (A) Collapse of viscosity data from Fig. 1 onto a universal curve assuming the avoided critical point form (KKZNT, black curve). Supplementary Figure 2 shows the residuals of the fit. (B) Data collapse of measurements reported in this work and for literature data for additional metallic glass-forming liquids. See Supplementary Table 2 for references to the data for the additional liquids broken down by composition.

All of these expressions fit the high-temperature viscosity data reasonably well, although most do not fit the entire temperature range. In particular, the most-commonly used VFT, and, to a lesser degree the MYEGA, expressions are both in poor agreement with the slope of the data near T_g (672K). Further, in the high temperature limit the MYEGA form approaches an Arrhenius temperature dependence far more weakly than predicted by the MD simulations⁴ and our experimental data. As a consequence, the fit values for η_0 are much larger than from fits to the other models and from the scaled data shown in Fig. 2B. The CG, DHTDSJ, BENK, and KKZNT models all fit the data over the entire temperature range. The DHTDSJ expression and to a lesser extent the CG form are, however, not consistent with a cross-over to a high temperature Arrhenius type behavior.

We now discuss a particular approximate expression to the universal function $F(z)$ appearing in our data collapse. In earlier studies, the avoided critical point expression (KKZNT) was shown to fit the viscosity data for many non-metallic liquids^{5–7}, albeit with five fitting parameters, compared to the three or four parameters for the other models investigated here. The expression may be written as $\eta = \eta_0 \exp(E/T)$ with a free energy barrier (expressed here in Kelvin) $E = E_\infty + T_A (bT_r)^z \Theta(T_A - T)$, where $\Theta(x)$ is the Heaviside function and the “reduced temperature” is $T_r \equiv (T_A - T)/T_A$. Earlier considerations suggested that some of those parameters are fixed. Bolstered by theory, empirical tests^{5–7} yielded an exponent $z \approx 8/3 \pm 1/3$. The KKZNT expression includes an “avoided critical point temperature” T^* , which like T_A in the MD results and the scaling temperature T_A introduced here, corresponds to the onset temperature for dynamical cooperativity. Based on theoretical considerations⁷, $T^*/T_l = 1.08$ for idealized liquids, where T_l is the liquidus, or melting, temperature—a tendency that is on average, is obeyed, but with significant spread. The universal curve further constrains the KKZNT expression, such that T_A and η_0 are the only remaining free parameters, with the values of the other parameters (now constants) determined by the fit to Vit 106a: $E_\infty = 6.466T_A$, $b = 4.536$, $z = 2.889$. As shown in Fig. 4A, with T_A and η_0 as the only free parameters this expression gives an excellent fit to all of the high temperature viscosity data shown in Fig. 1. The values for T_A and η_0 obtained from the fits are listed in Supplementary Table 2. The residuals of the fit are also plotted in Supplementary Figure 2. These T_A values are roughly 10% higher than those determined manually, with some scatter of a few percent. That the human eye would pick a lower temperature where the curvature is more apparent is unsurprising; the scatter is due to the difficulty in picking a temperature by eye in some alloys (e.g., Zr-Pt). These fit values are predictable from those obtained from the empirical scaling procedure used to find the universal curve for the high temperature data (Fig. 2B). Additionally, as will be demonstrated shortly, $T_A \approx 2.02T_g$ and (with less precision) $\eta_0 \sim nh$. Thus, the KKZNT fits to the universal collapse are nearly parameter free. As shown in Fig. 4B, this same constrained expression also fits data obtained by other investigators^{17,23–33} over a wide range of metallic glass families. Excluding a section for which measurements cannot be currently made, the universal curve fits well over 16 orders of magnitude in the viscosity. The reason for the small deviation in the high temperature data for Vit 1¹⁷ is likely due to a reported fragility transition in that liquid³⁴; no similar deviation was observed in our measured data for other liquids. Such good agreement for a range of different metallic liquids provides a striking demonstration of the validity of the KKZNT expression as an approximate functional form for the universal curve. Nevertheless, the possibility of other reasonable approximate forms is not ruled out. For example, the BENK expression (Table 1) also gives good agreement. The precise functional form for $\eta = \eta_0 F(T/T_A)$ is a matter for future theoretical studies. However, the KKZNT gives sufficiently good agreement with the experimental data over a very wide

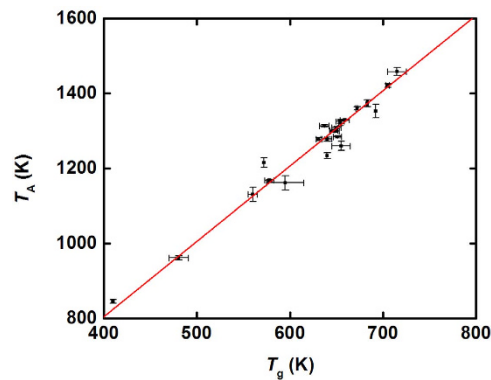


Figure 5. Comparison between the experimentally measured glass transition temperature, T_g , and T_A from the fits to the high temperature viscosity data. The solid line is a fit to the data, giving $T_A = (2.02 \pm 0.015)T_g$. Error bars are one s.d.

temperature range that it can be used here to examine the nature of the experimental scaling parameters.

Figure 5 shows the correlation between the values of T_A and T_g for the glass-forming liquids. The T_g values were obtained by us and by others from differential scanning calorimetry measurements, using a range of heating rates from 10 to 40 K per minute. The fit line shows that $T_A/T_g = 2.02 \pm 0.015$. This correlation is remarkable, and suggests a deep connection between the onset of cooperative dynamics in the liquid and the dynamical slowing down at the glass transition temperature. It is also the reason that T_g remains a useful scaling temperature.

This correlation also suggests novel approaches for the search for good glass forming liquids, at least for metallic glasses. For example, assuming the Turnbull criterion of good glass formability when (T_g/T_l) is large³⁵, makes it possible to assess trends in glass formability from liquid data alone, without actually forming a glass and measuring T_g . A study of the change in T_A values with the chemical composition of the liquid would show whether such a survey is practically useful.

Further Discussion of the Scaling Parameters. As discussed, the scaling parameters η_0 and T_A can be obtained from fits to the KKZNT theory. However, it is important to underscore that they can also be determined empirically by collapsing all of the experimental data onto a universal curve, making them fundamentally theory independent. Several interesting points emerge from an examination of the values obtained for the parameters.

The values for the extrapolated high-temperature viscosity, η_0 , suggest that there may exist a universal high-temperature limit of the viscosity. For the liquid metals studied here, this is *on average* equal to $n\hbar$ ($\langle \eta_0 \rangle = n\hbar$)¹², where \hbar is Planck's constant and n is the particle number per unit volume and the brackets $\langle \rangle$ denote an average over all of the liquids studied. Such typical values for η_0 have been predicted^{11,12} and may have an even deeper significance that extends beyond that of liquid metals. Fundamental lower limits on the viscosity are discussed widely in various contexts, with recent interest³⁶ driven by predictions from string theory and holographic dualities, which were compared with measurements at the Relativistic Heavy Ion Collider³⁷.

The strong correlation between T_A and T_g ($T_A \sim 2.02 T_g$) for all of the metallic-glass-forming liquids examined supports a long-held belief from other complementary approaches that glass formation might be a consequence of a high temperature transition crossover. Theories of this crossover include an avoided critical point⁵⁻⁷, a random first order transition^{38,39}, and mode coupling theories⁴⁰, among others (e.g.⁴¹). In the avoided critical point theory T_A corresponds to the transition temperature of the supercooled liquid in an idealized template – a transition that is avoided by frustration. There have been previous experimental hints of non-trivial dynamics associated with a viable dynamical crossover temperature, T_{cross} (above T_g). These include (a) the appearance (at $T < T_{\text{cross}}$) of short time, or β , relaxation processes accompanying the primary, or α , relaxation rates that are the focus of this work⁴², (b) the broadening of relaxation times about these two principal processes (typically this broadening is manifest in response functions that have a stretched exponential behavior)^{20,43}, (c) nonuniform dynamics in different spatial regions (dynamical heterogeneities)⁴⁴, (d) violation of the Stokes-Einstein relation⁴⁵, and (e) decoupling of translational and rotational diffusivity⁴⁶, and (f) phonon localization⁴. These phenomena appear and are strongly indicative of transformations that have an onset above T_g , yet at temperatures that are lower than $2 T_g$ in most studied non-metallic liquid systems. It is also important to note that T_A is much above predicted dynamic crossover temperatures⁴¹ and the mode-coupling temperature⁴⁷.

Emerging Questions. The results presented here raise several questions. For example, what is the origin of the observed universal behavior and the connection between T_g and T_A ? It has long been

known that many supercooled metallic liquids veer towards locally preferred low energy icosahedral structures^{5–7,48–50}, a tendency that other liquids generally do not share. This general propensity towards locally preferred structures lies at the origin of the avoided critical point model^{5–7}. MD simulations show that on decreasing the temperature below T_A many metallic glass forming liquids progressively develop more pronounced icosahedral order, with a length scale of connected icosahedral cluster networks that monotonically increases until they percolate throughout the entire system near T_g (see, e.g., Figure 7 of Ref. 10). This is in agreement with predications from avoided critical point theory^{5–7}, and could be the source of the connection between T_g and T_A that is experimentally observed here. Additionally, over the ensemble of metallic liquids that were examined, $T_A/T_1 \approx 1.075 \pm 0.188$ (where T_1 is the liquidus temperature), consistent with estimates suggested by the avoided critical theory⁷. However, it should be emphasized that local order need not be icosahedral for theories of an avoided critical point nature to be valid. The local structures of Pd-Si, for example, are likely not strongly icosahedral, while some liquids that have more definitive icosahedral order and still fit the universal curve (e.g., Zr-Pt) deviate more from the predictions of the scaling parameters η_0 and T_A than do others.

A second question is what these results imply about liquid fragility. Metallic liquids show a range of fragility, but occupy the central part of an Angell plot¹. The stronger metallic liquids (e.g. Vit106) are more fragile than strong liquids like SiO₂. If η_0 were truly independent of temperature for liquid metals, as suggested from the data presented, and if T_A/T_g were truly constant, then the fragility index, defined as $m \equiv (\partial \log_{10} \eta / \partial (T_g/T))_{T=T_g}$, would be the same for all metallic liquids. This is extremely unlikely, however. Evidence for fragility exists not only in the dynamical properties (where typically an assumed VFT form for η is invoked), but also in thermodynamic properties^{51,52} and rate of structural ordering^{3,53}. Our studies show that fragility is embedded in the nature of the deviation of the T_A/T_g ratio found for different metallic liquids as well as possible remnant small deviations of the viscosity data from a universal collapse. It is useful to see how the fragility index m varies with T_A/T_g for all of the liquids studied. For metallic liquids fragility is frequently determined from calorimetric measurements⁵⁴ in addition to or when viscosity data are unobtainable. However, while this gives reliable values of m for good glass-formers, the values for marginal glass forming liquids are scarce and often unreliable. This makes it impossible at present to examine the correlation for all of the liquids studied. Instead, Supplementary Fig. 1 compares the ratio T_A/T_g with values of m reported^{13,55–57} for the good glass-forming liquids studied here with. The m values range from about 32 to 75; for comparison the m values for SiO₂ and o-terphenyl are 20 and 81, respectively. While there are substantial disparities in the values of m that different groups have reported for any single metallic liquid, the average reported fragility index values clearly increase with T_A/T_g , in agreement with a structural origin of fragility (see Ref. 53). However, as seen from the steepness of the slope in Figure S1, the large variations in the values of m evaluated just above T_g are not similarly reflected in the T_A/T_g ratio. The origin of this enigmatic behavior is not understood.

Summary and Conclusions

In summary, the measured data suggest a new universality in the dynamical behavior of liquid metals when the temperature is scaled by T_A , which corresponds to the onset of dynamical cooperativity, and when the extrapolated infinite temperature viscosities (η_0) are properly accounted for. That the glass transition temperature, T_g , is strongly correlated with T_A suggests that the cooperative processes that eventually lead to the glass transition are already present in the high temperature liquid. The rapid cooling rates needed for molecular dynamics (MD) studies often raise questions of the validity of comparisons between computed results and experimental data for temperatures near T_g . The results presented here, however, show that it is possible to more realistically probe processes associated with the glass transition by comparing computed results with experimental data obtained at high-temperature, where experimental and MD relaxation times are comparable. It remains to be seen whether the results and conclusions presented here extend to liquids other than metallic ones, but our data for the metallic liquids clearly motivate the need for further investigation.

References

1. Angell, C. A. Formation of Glasses from Liquids and Biopolymers. *Science* **267**, 1924–1935, doi: 10.1126/Science.267.5206.1924 (1995).
2. Bendert, J. C. & Kelton, K. F. Correlation between kinetic strength, volumetric properties, and glass forming ability in metallic liquids. *Journal of Non-Crystalline Solids* **376**, 205–208, doi: 10.1016/J.jnoncrysol.2013.05.032 (2013).
3. Mauro, N. A., Vogt, A. J., Johnson, M. L., Bendert, J. C. & Kelton, K. F. Anomalous structural evolution in Cu₅₀Zr₅₀ glass-forming liquids. *Applied Physics Letters* **103**, 021904–021904–021904, doi: 10.1063/1.4813389 (2013).
4. Iwashita, T., Nicholson, D. M. & Egami, T. Elementary Excitations and Crossover Phenomenon in Liquids. *Physical Review Letters* **110**, 205504, doi: 10.1103/PhysRevlett.110.205504 (2013).
5. Kivelson, D., Kivelson, S. A., Zhao, X., Nussinov, Z. & Tarjus, G. A thermodynamic theory of supercooled liquids. *Physica A: Statistical Mechanics and its Applications* **219**, 27–38, doi: 10.1016/0378-4371(95)00140-3 (1995).
6. Tarjus, G., Kivelson, S. A., Nussinov, Z. & Viot, P. The frustration-based approach of supercooled liquids and the glass transition: a review and critical assessment. *Journal of Physics: Condensed Matter* **17**, R1143–R1182, doi: 10.1088/0953-8984/17/50/R01 (2005).
7. Nussinov, Z. Avoided phase transitions and glassy dynamics in geometrically frustrated systems and non-Abelian theories. *Physical Review B* **69**, 014208, doi: 10.1103/PhysRevB.69.014208 (2004).

8. Kivelson, D., Tarjus, G., Zhao, X. & Kivelson, S. A. Fitting of viscosity: Distinguishing the temperature dependences predicted by various models of supercooled liquids. *Physical Review E* **53**, 751 (1996).
9. Tarjus, G., Kivelson, D., Mossa, S. & Alba-Simionesco, C. Disentangling density and temperature effects in the viscous slowing down of glass-forming liquids. *Journal of Chemical Physics* **120**, 6135–6141, doi: 10.1063/1.1649732 (2004).
10. Soklaski, R., Nussinov, Z., Markow, Z., Kelton, K. F. & Yang, L. Connectivity of icosahedral network and a dramatically growing static length scale in Cu-Zr binary metallic glasses. *Physical Review B* **87**, 184203, doi: 10.1103/PhysRevB.87.184203 (2013).
11. Eyring, H. Viscosity, Plasticity, and Diffusion as Examples of Absolute Reaction Rates. *The Journal of chemical physics* **4**, 283, doi: 10.1063/1.1749836 (1936).
12. Nussinov, Z., Nogueira, F., Blodgett, M. & Kelton, K. F. Possible sharp quantum relaxation times in “classical” fluids: theory and experiment. *ArXiv preprint*, arXiv: **1409**, 1915 (2014). <http://adsabs.harvard.edu/abs/2014arXiv1409.1915N%3E.
13. Evenson, Z., Raedersdorf, S., Gallino, I. & Busch, R. Equilibrium viscosity of Zr–Cu–Ni–Al–Nb bulk metallic glasses. *Scripta Materialia* **63**, 573–576, doi: 10.1016/j.scriptamat.2010.06.008 (2010).
14. Rault, J. Origin of the Vogel–Fulcher–Tammann law in glass-forming materials: the α – β bifurcation. *Journal of Non-Crystalline Solids* **271**, 177–217 (2000).
15. Mauro, J. C., Yue, Y., Ellison, A. J., Gupta, P. K. & Allan, D. C. Viscosity of glass-forming liquids. *Proceedings of the National Academy of Sciences of the United States of America* **106**, 19780–19784, doi: 10.1073/pnas.0911705106 (2009).
16. Cohen, M. H. & Grest, G. S. Liquid-Glass Transition, a Free-Volume Approach. *Physical Review B* **20**, 1077–1098, doi: 10.1103/PhysRevB.20.1077 (1979).
17. Demetriou, M. D. *et al.* Cooperative shear model for the rheology of glass-forming metallic liquids. *Physical Review Letters* **97**, 065502, doi: 10.1103/PhysRevLett.97.065502 (2006).
18. Elmatad, Y. S., Chandler, D. & Garrahan, J. P. Corresponding states of structural glass formers. *The Journal of Physical Chemistry B* **113**, 5563–5567, doi: 10.1021/jp810362g (2009).
19. Elmatad, Y. S., Jack, R. L., Chandler, D. & Garrahan, J. P. Finite-temperature critical point of a glass transition. *Proceedings of the National Academy of Sciences of the United States of America* **107**, 12793–12798, doi: 10.1073/pnas.1006306107 (2010).
20. Mallamace, F. *et al.* Transport properties of glass-forming liquids suggest that dynamic crossover temperature is as important as the glass transition temperature. *Proceedings of the National Academy of Sciences of the United States of America* **107**, 22457–22462, doi: 10.1073/Pnas.1015340107 (2010).
21. Rossler, E., Hess, K. U. & Novikov, V. N. Universal representation of viscosity in glass forming liquids. *Journal of Non-Crystalline Solids* **223**, 207–222, doi: 10.1016/S0022-3093(97)00365-7 (1998).
22. Schmidtke, B., Petzold, N., Kahlau, R., Hofmann, M. & Rossler, E. A. From boiling point to glass transition temperature: Transport coefficients in molecular liquids follow three-parameter scaling. *Physical Review E* **86**, doi: 10.1103/PhysRevE.86.041507 (2012).
23. Kawamura, Y., Nakamura, T., Kato, H., Mano, H. & Inoue, A. Newtonian and non-Newtonian viscosity of supercooled liquid in metallic glasses. *Materials Science & Engineering A* **304**, 674–678, doi: 10.1016/S0921-5093(00)01562-8 (2001).
24. Tsang, K. H., Lee, S. K. & Kui, H. W. Viscosity of molten Pd₄₀Ni₄₀P₂₀. *J Appl Phys* **70**, 4837–4841, doi: 10.1063/1.349050 (1991).
25. Tsang, K. H. & Kui, H. W. Viscosity of molten Pd₈₂Si₁₈ and the scaling of viscosities of glass forming systems. *Journal of Applied Physics* **72**, 93–96, doi: 10.1063/1.352101 (1992).
26. Stojanova, L., Russev, K. & Illekova, E. Study of the Structural Relaxation of Pd₈₂Si₁₈ Metallic-Glass by Thermal-Expansion and Viscous-Flow Measurements. *Materials Science & Engineering A* **133**, 529–531, doi: 10.1016/0921-5093(91)90126-8 (1991).
27. Hyers, R. *et al.* Surface tension and viscosity of quasicrystal-forming Ti–Zr–Ni alloys. *International Journal of Thermophysics* **25**, 1155–1162 (2004).
28. Bradshaw, R. C. *et al.* Containerless Measurements of Thermophysical Properties of Zr₅₄Ti₈Cu₂₀Al₁₀Ni₈. *Annals of the New York Academy of Sciences* **1077**, 63–74, doi: 10.1196/annals.1362.058 (2006).
29. Bradshaw, R. C. *et al.* Nonlinearities in the undercooled properties of Ti_{39.5}Zr_{39.5}Ni₂₁. *Philosophical Magazine* **86**, 341–347, doi: 10.1080/14786430500253968 (2006).
30. Wu, S.-S., Chin, T.-S., Su, K.-C. & Shyr, F.-S. Undercooled Liquid Viscosity and Glass Formation of a Mg₆₂Cu₂₆Y₁₂Alloy. *Japanese Journal of Applied Physics* **35**, 175–178, doi: 10.1143/JJAP.35.175 (1996).
31. Busch, R., Liu, W. & Johnson, W. L. Thermodynamics and kinetics of the Mg₆₅Cu₂₅Y₁₀ bulk metallic glass forming liquid. *Journal of Applied Physics* **83**, 4134, doi: 10.1063/1.367167 (1998).
32. Haumesser, P.-H., Bancillon, J., Daniel, M., Perez, M. & Garandet, J.-P. High-temperature contactless viscosity measurements by the gas–film levitation technique: Application to oxide and metallic glasses. *Review of Scientific Instruments* **73**, 3275–3285, doi: 10.1063/1.1499756 (2002).
33. Lu, I. R., Görler, G. P., Fecht, H. J. & Willnecker, R. Investigation of specific volume of glass-forming Pd–Cu–Ni–P alloy in the liquid, vitreous and crystalline state. *Journal of Non-Crystalline Solids* **312–314**, 547–551, doi: 10.1016/S0022-3093(02)01767-2 (2002).
34. Way, W. & Busch, R. The influence of shear rate and temperature on the viscosity and fragility of the Zr_{41.2}Ti_{13.8}Cu_{12.5}Ni_{10.0}Be_{22.5} metallic-glass-forming liquid. *Acta Materialia* **55**, 2977–2983, doi: 10.1016/j.actamat.2006.12.032 (2007).
35. Turnbull, D. Under what conditions can a glass be formed? *Contemporary Physics* **10**, 473–488, doi: 10.1080/00107516908204405 (1969).
36. Kovtun, P. K., Son, D. T. & Starinets, A. O. Viscosity in strongly interacting quantum field theories from black hole physics. *Physical Review Letters* **94**, 111601 (2005).
37. Shuryak, E. Why does the quark–gluon plasma at RHIC behave as a nearly ideal fluid? *Progress in Particle and Nuclear Physics* **53**, 273–303, doi: 10.1016/j.ppnp.2004.02.025 (2004).
38. Kirkpatrick, T., Thirumalai, D. & Wolynes, P. Scaling concepts for the dynamics of viscous liquids near an ideal glassy state. *Physical Review A* **40**, 1045–1054, doi: 10.1103/PhysRevA.40.1045 (1989).
39. Lubchenko, V. & Wolynes, P. G. Theory of structural glasses and supercooled liquids. *Annual Review of Physical Chemistry* **58**, 235–266, doi: 10.1146/annurev.physchem.58.032806.104653 (2007).
40. Götze, W. Recent tests of the mode-coupling theory for glassy dynamics. *Journal of Physics: Condensed Matter* **11**, A1–A45, doi: 10.1088/0953-8984/11/10A/002 (1999).
41. Schröder, T. B., Sastry, S., Dyre, J. C. & Glotzer, S. C. Crossover to potential energy landscape dominated dynamics in a model glass-forming liquid. *Journal of Chemical Physics* **112**, 9834–9840, doi: 10.1063/1.481621 (2000).
42. Mallamace, F. *et al.* On the ergodicity of supercooled molecular glass-forming liquids at the dynamical arrest: the o-terphenyl case. *Scientific Reports* **4**, 3747, doi: 10.1038/srep03747 (2014).
43. van Megen, W., Mortensen, T., Williams, S. & Müller, J. Measurement of the self-intermediate scattering function of suspensions of hard spherical particles near the glass transition. *Physical Review E* **58**, 6073–6085, doi: 10.1103/PhysRevE.58.6073 (1998).
44. Ediger, M. D. Spatially heterogeneous dynamics in supercooled liquids. *Annual Review of Physical Chemistry* **51**, 99–128, doi: 10.1146/annurev.physchem.51.1.99 (2000).
45. Tarjus, G. & Kivelson, D. Breakdown of the Stokes–Einstein relation in supercooled liquids. *Journal of Chemical Physics* **103**, 3071, doi: 10.1063/1.470495 (1995).

46. Fujara, F., Geil, B., Sillescu, H. & Fleischer, G. Translational and Rotational Diffusion in Supercooled Orthoterphenyl Close to the Glass-Transition. *Zeitschrift für Physik B Condensed Matter* **88**, 195–204, doi: 10.1007/Bf01323572 (1992).
47. Götze, W. *Complex Dynamics of Glass-Forming Liquids: A Mode-Coupling Theory*. (OUP Oxford, 2008).
48. Kelton, K. F. *et al.* First x-ray scattering studies on electrostatically levitated metallic liquids: demonstrated influence of local icosahedral order on the nucleation barrier. *Physical Review Letters* **90**, 195504, doi: 10.1103/PhysRevLett.90.195504 (2003).
49. Schenk, T., Holland-Moritz, D., Simonet, V., Bellissent, R. & Herlach, D. M. Icosahedral short-range order in deeply undercooled metallic melts. *Physical Review Letters* **89**, 075507 (2002).
50. Frank, F. C. Supercooling of Liquids. *Proceedings of the Royal Society of London. Series A. Mathematical and Physical Sciences* **215**, 43–46, doi: 10.1098/rspa.1952.0194 (1952).
51. Ito, K., Moynihan, C. T. & Angell, C. A. Thermodynamic determination of fragility in liquids and a fragile-to-strong liquid transition in water. *Nature* **398**, 492–495 (1999).
52. Martinez, L. M. & Angell, C. A. A thermodynamic connection to the fragility of glass-forming liquids. *Nature* **410**, 663–667, doi: 10.1038/35070517 (2001).
53. Mauro, N. A., Blodgett, M., Johnson, M. L., Vogt, A. J. & Kelton, K. F. A structural signature of liquid fragility. *Nature Communications* **5**, doi: 10.1038/ncomms5616 (2014).
54. Evenson, Z., Gallino, I. & Busch, R. The effect of cooling rates on the apparent fragility of Zr-based bulk metallic glasses. *Journal of Applied Physics* **107**, doi: 10.1063/1.3452381 (2010).
55. Perera, D. N. Compilation of the fragility parameters for several glass-forming metallic alloys. *Journal of Physics: Condensed Matter* **11**, 3807–3812, doi: 10.1088/0953-8984/11/19/303 (1999).
56. Johnson, W. L., Demetriou, M. D., Harmon, J. S., Lind, M. L. & Samwer, K. Rheology and ultrasonic properties of metallic glass-forming liquids: A potential energy landscape perspective. *MRS Bulletin* **32**, 644–650 (2007).
57. Takeuchi, A., Kato, H. & Inoue, A. Vogel-Fulcher-Tammann plot for viscosity scaled with temperature interval between actual and ideal glass transitions for metallic glasses in liquid and supercooled liquid states. *Intermetallics* **18**, 406–411, doi: 10.1016/j.intermet.2009.08.015 (2010).

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Author Contributions

K.K. supervised the project, coordinated the experimental and theoretical analysis, and prepared the manuscript, in collaboration with the other authors. He, in collaboration with T.E. conceived of and demonstrated the possibility of the universal curve. M.B. made the experimental measurements of viscosity. The more detailed analysis of the experimental data was mainly carried out by M.B. with suggestions from Z.N. and K.K. Z.N. and T.E. provided theoretical support. Z.N. examined viable physical content/consequences of the scaling collapse parameters and fits.

Additional Information

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