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Universal structural softening in metallic glasses indicated by boson heat capacity peak


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Low-temperature heat capacity is systematically investigated in various glassy and crystalline polymorphs of a wide range of metallic glasses. We reveal that the boson heat capacity peak beyond the Debye level arises from both excess phonon scattering and background electronic excitation, and the two contributions are strongly coupled and also material-dependent. It is interesting to observe that the boson heat capacity peaks obey an inversely linear correlation between their heights and characteristic positions, which is mainly dominated by phononic anomalies. This indicates a universal structural softening among the studied glasses when the boson peak occurs. We further suggest a possibility that the linear evolution of the fast boson peaks can probe into the slow structural softening across the glass transition, and the two dynamic processes are controlled by the short-time shear modulus associated with local soft regions in fragile glasses. Published by AIP Publishing. https://doi.org/10.1063/1.5016984

The nature of vibrations in glasses belongs to the grand mysteries in condensed matter physics. At terahertz (THz) frequencies, glasses generally exhibit an anomaly in their vibrational spectrum, i.e., the vibrational density of states (VDOSSs) exceeds over the Debye level determined by continuum mechanics. This enhancement of THz excitations in glasses is called the “boson peak (BP)”. It is commonly believed that the BP pertains to all glasses, irrespective of the bond types or the compositions and thus also including metallic glasses. Despite much controversy, it now seems accepted that the BP originates from transverse phonons mixed with localized soft modes. The extra scattering of quasilocalized phonons contributes to an excess in the low-temperature \( T \) heat capacity \( C_p \) as compared to the Debye \( T^3 \) law. Thus, the excess heat capacity occurs visibly as a peak (namely, the “boson heat capacity peak”) in the plot of \( C_p/T^3 \) vs \( T \) at temperatures of \( \sim 10 \text{ K} \) for all glasses.

Although it has not yet been completely understood, the BP opens a window into probing the intrinsic structural heterogeneity that plays a crucial role in glass relaxation and flow. Some intriguing correlations have been discovered between fast BP excitations and slow structural dynamics in glasses. For example, Sokolov et al. found that the BP intensity increases with a decrease in the kinetic fragility. Experiments show that either mechanical deformation or isothermal aging can induce the evolution of the BP height and position. Recent simulations clearly demonstrate that quasilocalized soft modes (contributing to BPs) correlate strongly with fertile regions for irreversible rearrangements or shear transformations.

Regardless of the specific reasons, the BP evolution itself exhibits a common feature: the stronger BPs always shift to lower frequencies/temperatures and vice versa. The universality of this behavior implies that the underlying origin of BPs may be quite fundamental in glasses albeit the diversity of microscopic constituents. Due to the structural simplicity and similarity, metallic glasses offer ideal model systems for exploring the evolution of the BPs. For metallic glasses, however, it is more complex to experimentally define the BP as a peak in \( C_p/T^3 \) vs \( T \). This is because the electronic heat capacity inevitably interferes at such low temperatures. At least the boson heat capacity peak has to include quasilocalized phonon and background electronic contributions, and sometimes, the latter could hide the former, due to a strong electron-phonon coupling. Naturally, one would like to ask how the boson heat capacity peak evolves in metallic glasses with the inherent interaction of electrons and phonons, which is not covered previously.

In this Letter, we report a systematic study on the boson heat capacity peaks of various glassy and crystalline polymorphs of a wide range of metallic glasses. By examining the Sommerfeld coefficient \( \gamma \), it is demonstrated that the electronic contribution to \( C_p \) is extremely sensitive to the topological and chemical disorder of glasses. The significant electronic contribution will hide the BPs in \( C_p/T^3 \)-plots, causing an apparent anomaly of their evolution. By subtracting the electronic part from the low-temperature \( C_p \), we can locate the boson peak whose evolution still exhibits the common

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feature. More importantly, it is revealed that the heights of the boson heat capacity peaks follow an inversely linear relationship with temperature at which \( C_p/T^3 \) is maximum. This implies a universal structural softening in metallic glasses when the BP occurs. The BP-induced softening shows a possible link to the structural softening across the glass transition, and both dynamic processes are shear-dominated on short timescales.3,19

Vitreloy 1, with a nominal composition of Zr_{41.2}Ti_{13.8}Cu_{12.5}Ni_{10}Be_{22.5} (at. %), was chosen as a model metallic glass to show the evolution of boson heat capacity peaks in different topological states. The low-temperature (1.9 K \( \leq T \leq 100 \) K) specific heat capacity \( C_p \) was measured using a Quantum Design physical property measurement system (PPMS) on the as-cast glassy, annealed glassy, and fully polycrystalline states of the Vitreloy 1. The supplementary material describes how the three Vitreloy 1 states have been synthesized and characterized.

Figure 1(a) plots the specific heat capacity \( C_p \) for the as-cast, annealed, and crystallized Vitreloy 1 in the temperature interval of 1.9–100 K. The inset shows the corresponding log-log plot. As expected, the heat capacity \( C_p \) for each state decreases with decreasing temperature. All \( C_p \) data will approach zero if \( T \rightarrow 0 \) K, and their high-temperature limit is about 0.416 J g\(^{-1}\)K\(^{-1}\), determined by the Dulong-Petit law. The heat treatment reduces the heat capacity by deepening inherent structures (ISs) of glasses.20,21 Figure 1(b) plots the specific heat capacity data as \( C_p/T^3 \) vs \( T \) to reveal the BPs beyond the Debye level. In this plot, the ordering-induced decrease of \( C_p \) becomes more apparent. For the as-cast and annealed Vitreloy 1 glasses, the BP only appears as an obscure shoulder and cannot be readily identified. However, the Vitreloy 1 polycrystal exhibits a visible BP-like anomaly at about 20 K. This kind of anomaly has been previously observed in crystalline materials,7,22,23 always showing a significantly higher peak temperature/frequency compared to that of the glass BPs. We believe that the currently observed \( C_p \) anomaly stems from disordered lattices at nanograin boundaries that abound in the polycrystalline alloys (see supplementary material Fig. S3). The underlying mechanism is the transverse acoustic (TA) van Hove singularity,17,23 a suggested counterpart of the glass BPs in crystals. A question thus arises why the Vitreloy 1 glass with a higher degree of disorder does not show a pronounced BP? According to the seminal study of Li et al.,24 such a paradox is attributed to the contribution of electronic excitations to low-temperature \( C_p \).

The electronic \( C_p \) is typical of metallic glasses or crystalline alloys but does not exist in traditional glasses without metallic bonding. It is thus speculate that the electronic \( C_p \) might be relatively weak in the alloys with higher structural order. In other words, the electronic \( C_p \) contribution should highly depend on the topological disorder/order in alloys.

We determine the electronic \( C_p \) contribution by plotting \( C_p/T \) vs \( T^2 \) below 8 K for the as-cast, annealed, and crystallized Vitreloy 1 (see supplementary material Fig. S4). The low-temperature \( C_p \) data can be well fitted with the formula \( C_p/T = \beta T^2 + \gamma \), where the Sommerfeld coefficient \( \gamma \) measures the degree of electronic contribution. It is worth noting that \( \gamma \) becomes smaller for the heavily heat treated Vitreloy 1. For the annealed states, \( \gamma \) decreases by about 3% as compared with that of the as-cast, and \( \gamma \) of the crystallized states decreases by about 33%. This behavior implies a strong electron-phonon coupling that contributes to the low-temperature \( C_p \) in glassy or crystalline states of the alloy. The electronic \( C_p \) contribution becomes more significant in highly disordered alloys. Contrarily, topological/structural ordering induced by heat treatment decreases the electronic contribution \( \gamma \), making the boson heat capacity peaks more visible. This explains why the polycrystalline Vitreloy 1 exhibits a pronounced “BP” that is a multiplex consequence of the plentiful disordered boundaries (TA singularity) and relatively weak electronic contribution to \( C_p \).

Our analysis confirms that the boson heat capacity peaks due to phononic anomalies can be determined by subtracting the linear electronic contribution \( \gamma T \). Actually, this approach has been widely adopted in metallic glasses.6,8,24,25 An alternative approach defines a boson peak as the difference between \( C_p/T^3 \) of the glass and the corresponding crystal. However, it is very difficult to find such a reference crystal that does not show a BP-like \( C_p \) anomaly and meanwhile has the same electronic \( C_p \) as that of the glass. Here, we adopt the first approach because we believe that it reflects the essential properties of BPs.

Figure 2 plots the specific heat capacity as \( C_p/T^3 \) and \( (C_p - \gamma T)/T^3 \) vs \( T \) for the as-cast, annealed, and crystallized
Vitreloy 1 (from left to right). As compared with $C_p = T^3$, the $\frac{C_p}{C_0} = T^3$ curves clearly show where the BPs take place, and the position temperatures $T_{BP}$ of the BPs are readily located. More importantly, we obtain two kinds of boson heat capacity peaks. One as a peak in the $\frac{C_p}{C_0} = T^3$ plot is only due to the atomic vibrational or phononic anomaly, denoted as $BP_{ph}$. The other involves the electronic contribution $C_T$ in addition to the phononic $BP_{ph}$, denoted as $BP_{ph+el}$ in the $C_p = T^3$ plot. The heights corresponding to $BP_{ph}$ and $BP_{ph+el}$ can be determined, respectively, as

$$H_{ph} = \frac{C_p \gamma T}{T^3} T_{BP},$$

$$H_{ph+el} = \frac{C_p T}{T^3} T_{BP}.$$  

For Vitreloy 1, the evolution of both $BP_{ph}$ and $BP_{ph+el}$ induced by structural changes still follows the common feature. The stronger (higher $H_{ph}$ or $H_{ph+el}$) the boson heat capacity peaks are the lower the corresponding temperatures ($T_{BP}$) are observed. The as-cast Vitreloy 1 glass that corresponds to the state of the highest disorder shows the strongest BPs where both phononic and electronic contributions are highest. The polycrystalline Vitreloy 1 shows the lowest “BPs” with the weakest phononic and electronic contributions. The BP of the annealed glassy Vitreloy 1 takes an intermediate level.

The universality of evolution of boson heat capacity peaks has been further studied by comparing various metallic glasses by plotting their $H_{ph}$ vs $1/T_{BP}$. Obviously, stronger $BP_{ph}$ with higher $H_{ph}$ corresponds to lower $T_{BP}$ and vice versa. Disordered glasses formed by fast cooling move to the upper right corner of Fig. 3(a), while those well annealed reside in the lower left corner, as well as their crystallized states. There appears to be a general linear trend between $H_{ph}$ and $T_{BP}$; however, the data are noted to a little bit scatter. We believe that this scatter results from the inherent effect of the additional electronic $C_p$ contribution. It is indeed found that the Sommerfeld coefficients $\gamma$ are extremely sensitive to glassy structures and compositions (see supplementary material Table S1). Such a strong electron-phonon coupling leads to the material-dependent contribution to BP from either phonons or electrons. Quite interestingly, when we plot $H_{ph+el}$ vs $1/T_{BP}$, as shown in Fig. 3(b), all data collapse well onto a single master straight line. This indicates that the combination of the two material-dependent contributions does not depend on specific glasses any longer. The observed linear correlation can be well fitted by

$$H_{ph+el} = \left(\frac{C_p}{T^3}\right)_{T_{BP}} = \Gamma \frac{1}{T_{BP}} + \Phi,$$  

Figure 3(a) presents the phononic $BP_{ph}$ evolution among various metallic glasses by plotting their $H_{ph}$ vs $1/T_{BP}$. Obviously, stronger $BP_{ph}$ with higher $H_{ph}$ corresponds to lower $T_{BP}$ and vice versa. Disordered glasses formed by fast cooling move to the upper right corner of Fig. 3(a), while those well annealed reside in the lower left corner, as well as their crystallized states. There appears to be a general linear trend between $H_{ph}$ and $T_{BP}$; however, the data are noted to a little bit scatter. We believe that this scatter results from the inherent effect of the additional electronic $C_p$ contribution. It is indeed found that the Sommerfeld coefficients $\gamma$ are extremely sensitive to glassy structures and compositions (see supplementary material Table S1). Such a strong electron-phonon coupling leads to the material-dependent contribution to BP from either phonons or electrons. Quite interestingly, when we plot $H_{ph+el}$ vs $1/T_{BP}$, as shown in Fig. 3(b), all data collapse well onto a single master straight line. This indicates that the combination of the two material-dependent contributions does not depend on specific glasses any longer. The observed linear correlation can be well fitted by

$$H_{ph+el} = \left(\frac{C_p}{T^3}\right)_{T_{BP}} = \Gamma \frac{1}{T_{BP}} + \Phi,$$  

FIG. 2. Low-temperature $C_p/T^3$ and $(C_p - \gamma T)/T^3$ vs $T$ for the as-cast, annealed, and crystallized Vitreloy 1 (from left to right).

FIG. 3. Evolution of boson capacity heat peaks in metallic glasses by plotting their heights (a) $H_{ph}$ and (b) $H_{ph+el}$ as a function of positions $1/T_{BP}$. The solid line in (b) is the least-squares fitting by Eq. (3). Colorful spheres: data in this work; black squares: data from the literature.
where the universal slope ($\Gamma = 28.58$) implies the existence of a fundamental behavior of various glassy structures. The intercept ($\Phi = 0.014$) is supposed to be less meaningful since it is much less than $H_{\text{ph}+\text{el}}$ by at least 2 orders of magnitude.

Figure 3 clearly demonstrates that, although it is improved by the electronic contribution, the observed linear correlation (3) is mainly dominated by the extra atomic vibrations. In what follows, we only consider the dominated phononic effect that is actually coupled with the secondary electronic contribution. Therefore, this implies that the VDOS BPs of glasses in general may also show a linear evolution between their heights and characteristic frequencies, which has been supported by simulations \cite{31} and experiments. \cite{32,33} For example, Grigera et al. \cite{29} computed the VDOS $g(\omega)$ of the IS of a fragile glass using the Swap Monte Carlo algorithm. By changing the temperatures, they found that the height $g(\omega_{\text{BP}})/\omega_{\text{BP}}^2$ and the position $\omega_{\text{BP}}$ of the BPs satisfy a scaling law, which is only controlled by the IS energy. At temperatures well below the mode-coupling temperature, this scaling law approximates to an inversely linear behavior (or linear if adopting $1/\omega_{\text{BP}}$ like $1/T_{\text{BP}}$ here). Recently, using nuclear inelastic scattering, Chumakov et al. \cite{17} measured the iron-partial VDOS of a sodium silicate glass at various pressures. The phenomenon that the shift of BPs is nearly linear between their heights and positions can also be noted. In the present work, of great interest is that the different boson heat capacity peaks of various metallic glasses display an identical linear evolution. Such a universal behavior implies that the characteristic structures responsible for BPs are of, to some extent, similarity or resemblance for the studied glasses. As already discussed, the BP per se arises from quasilocalized soft modes, i.e., transverse vibrational modes associated with local soft regions (LSRs).\cite{3,13,14} The long-range elastic correlations of these LSRs cause a strong excess phonon scattering \cite{31} so that transverse acoustic phonons eventually lose their wave-like character, appearing as the BP. We note that the metallic glasses listed in supplementary material Table S1 belong to fragile liquids. In these glasses, LSRs construct a majority matrix phase surrounding strongly bonded regions. Therefore, the global shear modulus of glasses is mainly controlled by the local shear modulus of these LSRs, obeying the mechanical series model.\cite{32,33} In this scenario, Shintani and Tanaka \cite{3} suggested that the BP height is inversely proportional to the shear modulus. Vasiliev et al. \cite{22} further validated this simple relation and found that it is indeed controlled by the defect concentration akin to the fraction of LSRs. By the same token, we reasonably assume that

$$H_{\text{ph}+\text{el}} = \left(\frac{C_p}{T^3}\right)_{\text{BP}} \propto \frac{1}{G_{\infty}},$$  

(4)

where $G_{\infty}$ is the instantaneous shear modulus of glasses due to the extremely low-temperature ($\sim 10$ K) or high-frequency (THz) condition. Upon substituting Eq. (4) into Eq. (3), we get the expression of the slope $\Gamma$

$$\Gamma = \frac{1}{dG_{\infty}/dT_{\text{BP}}},$$  

(5)

It reveals that $1/\Gamma$ corresponds to the linear change of $G_{\infty}$ among various metallic glasses ($T_{\text{BP}}$), that is, $dG_{\infty}/dT_{\text{BP}} \approx 0.035 \text{ GPa/K}$; here, all dimension anharmonicities are observed in Eq. (4). This linear $G_{\infty}$-change actually reflects a universal sample-to-sample structural softening, which can be precisely probed by the shift of the boson capacity heat peaks.

In glass science, an idea that the instantaneous shear modulus $G_{\infty}$ can determine the glass flow is gaining increasing support from experiments,\cite{34} simulations\cite{35} to theories.\cite{36,19} A typical $G_{\infty}$-governed flow is the glass transition upon heating. As illustrated in Fig. 4, the glass transition can be represented as the $\text{z}$-relaxation hopping from a “metabasin” to a neighboring relatively shallower one in ISs of the potential energy landscape (PEL). Such a hopping event is essentially indistinguishable in the perspective of the static structures but can be detected by the decrease in $G_{\infty}$ that is the second derivative of the PEL of “metabasin”. As discussed above, the change of $G_{\infty}$ closely links with the evolution of BPs. It is therefore tempting to relate the $G_{\infty}$-governed softening at $T_{\text{BP}}$ to that at the glass transition temperature $T_g$, leading to

$$\Gamma = \frac{1}{dG_{\infty}/dT_{\text{BP}}} \approx \frac{1}{dG_{\infty}/dT_{T_g}}.$$  

(6)

To prove this speculation, data on $|dG_{\infty}/dT|_{T_g}$ are required. It is very difficult to precisely measure the instantaneous shear modulus $G_{\infty}$ of a frozen IS structure that undergoes the glass transition. Nevertheless, we identified three reports on experimental work, where the ultrasonic method was adopted to approximately frozen glass configurations. Lind et al.\cite{37} found that for the Vitreloy 4 (Zr46.75Ti8.25Cu7.5Ni10Be27.5) having different configurational states, their equilibrium liquid shows a strong linear relation $|dG_{\infty}/dT|_{T_g} = 0.035 \text{ GPa/K}$ around $T_g$. Keryvin et al.\cite{38} measured the change in Young’s modulus $E_{\infty}$ during a linear heating for a Zr55Cu34Ni9Al10 metallic glass. They found that there is a fast softening rate, $|dE_{\infty}/dT|_{T_g} = 0.089 \text{ GPa/K}$, at $T_g$. For metallic glasses, $G/E = 0.39$ is well satisfied.\cite{39} This also leads to $|dG_{\infty}/dT|_{T_g} \approx 0.035 \text{ GPa/K}$. Konnik et al.\cite{27} performed direct in situ
measurements of $G_\infty$ of the as-cast and annealed bulk Pd$_{40}$Cu$_{30}$Ni$_{10}$P$_{20}$ below and above $T_g$. A linear relation $|dG_\infty/dT|_P \approx 0.034\text{ GPa/K}$ was again observed. Such a striking agreement indicates a possibility that the THz BP anomaly and the glass transition on long timescales share the identical dynamic or even structural origin. Our results demonstrate that the sample-to-sample structural softening when the BP occurs is physically akin to the sample’s hopping on different “metabasins” in the ISs across the glass transition (see Fig. 4).

In conclusion, we systematically measured or analyzed the low-temperature heat capacity of various glassy and crystalline polymorphs of a wide range of metallic glasses. By distinguishing between the excess phononic scattering and background electronic excitation, we exactly locate the boson heat capacity peak. A main finding of the present study is that the boson heat capacity peaks involving both phononic and electronic contributions show an inversely linear correlation between their heights and position temperatures. The observed correlation implies a universal shear or structural softening among the studied metallic glasses, which can be understood in terms of transverse soft modes associated with local soft regions. We further bridge a possible link between the fast boson peak softening and the slow structural softening across the glass transition via the short-time shear modulus. Our work suggests that, mediated by $G_\infty$, the BP can be adopted to probe the glass transition. It must be pointed out that the present findings may not be valid for metallic glasses with significant magnetic heat capacity, heavy-fermion behavior, or belonging to strong liquids. In addition, for metallic glasses, the quantitative phonon-electron interaction deserves further investigations for deepening our fundamental understanding of low-temperature calorimetry anomalies.

See supplementary material for the preparation, heat treatments, and structural characterization of the Vitreloy 1 as well as the analytical data about the low-temperature heat capacity for various metallic glasses and their crystalline counterparts.

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