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ABSTRACT

Thermomechanical processing of metallic glasses can, remarkably, induce significant changes in structure and properties, even when the imposed macroscopic strains are well within the elastic regime. The changes can take the glasses to higher-energy “rejuvenated” or lower-energy “aged” states, with rejuvenation being of particular interest as it improves their mechanical properties. It has generally been assumed that the induced property changes would evolve monotonically with the extent of processing. We show that with sufficiently intense ultrasonic elastic processing, the intrinsic structural competition between damage and repair facilitated by increased atomic mobility can lead to oscillatory energy storage. The uncovering of this behavior forces reconsideration about the range of energy states attainable in metallic glasses by elastic deformation and may provide opportunities.

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Metallic glasses (MGs) are out-of-equilibrium systems with the dense packing structure of atoms but no long-range periodicity. MGs are characterized by yield stresses and elastic strain limits far exceeding those of conventional polycrystalline alloys, yet they show poor plasticity in compression due to strain softening and consequent localized flow in shear bands. For bulk MGs, strain softening is the principal hindrance to their wider application as structural materials. To improve the mechanical properties, many approaches have been adopted, from imposed patterns of residual stress to component size reduction. Thermomechanical processing (TMP) is established as a means for improving the mechanical functionality of metallic materials. When a polycrystalline metal is plastically deformed at room temperature, part of the mechanical work done is stored in the material through an increase in the concentration of defects such as dislocations, eventually leading to nanostructured and ultrafine-grained structures with high strength. Comparably, MGs also can store a fraction of applied cold work raising their internal energy to resemble states attainable by quenching the initial melt at higher rates. This rejuvenation leads to decreased yield stress and possibly improved plasticity. Plastic deformation can also induce more relaxed or aged states in MGs. Processing can therefore induce rejuvenation or aging. More intense processing, a lower processing temperature, and a lower-energy initial state favor rejuvenation.

In contrast to crystalline metals, MGs can even change their structure, gaining or losing energy, as a result of TMP in their nominally elastic regime. Although the property changes, rejuvenation, or aging induced in the elastic regime are usually monotonic, they can be more complex. Elastostatic compression first induces aging and then rejuvenation as the stress is increased, and under continued thermal cycling, the stored energy increases to a maximum and then decreases. In this study, we found that the ultrasonic elastic processing can induce an intrinsic structural competition between damage and repair and lead to oscillatory energy storage.
The amorphicity of the as-cast and UH-processed specimens was characterized by x-ray diffraction (XRD, a Bruker D8 instrument with Cu-Kα radiation). The morphology of treated specimens was examined by scanning electron microscopy (SEM). A power-compensated Perkin-Elmer DSC 8000 instrument, with a constant flow of high-purity argon, was used to investigate the structural relaxation behavior after the UH process. A heating rate of 0.33 K s⁻¹ and a cooling rate of 1.66 K s⁻¹ were used in all cases. During each DSC run, the specimen was first heated up to a fully crystallized state and cooled down to room temperature. Then, leaving the specimen in situ, a second heating scan was performed, the signal from the fully crystallized sample serving as the baseline correction when subtracted from the signal in the first scan. To account for some instrumental drift between scans, the measured signals were adjusted by a linear baseline correction to match the apparent specific heats in the supercooled liquid regions, as applied elsewhere.

The hardness measurements were carried out using a Vickers microhardness tester. Furthermore, cubic specimens of annealed Vit. 1 MG were treated by the UH process and then characterized by resonant ultrasound spectroscopy (RUS) and compression tests (see the supplementary material).

With static pressures in the ranges used (σ₀ = 3–96 MPa), the treated samples show no evidence of plastic deformation. Figures 1(d) and 1(e) show a disk of the commercial MG Vit. 1 after UH for 1.0 s at σ₀ = 96 MPa as a representative sample. The XRD pattern [inset in Fig. 1(d)] confirms that the struck surface of UH-treated samples remains amorphous. The temperature history [see the inset in Fig. 1(e)] shows that there is local heating, and the temperature of the struck surface increases up to 430 K. From this temperature profile, an atomic diffusion distance can be estimated, which is so small that any annealing effects of the temperature rise can be negligible. Furthermore, increasing σ₀ even to 1.44 GPa (corresponding to ~85% of the yield stress) leaves Vit. 1 samples undeformed, but larger σ₀ indeed ultimately fractures the sample and leads to remarkable shear banding throughout the disk (see Fig. S1, supplementary material).

The effects of UH on the as-cast specimens were monitored using DSC. Figure 2(a) shows the broad spectra of exothermic relaxations preceding glass transition temperature Tg for the as-cast specimens (DSC traces for the whole temperature range are shown in Fig. S2). The enthalpy of relaxation ΔHrel can be calculated by integrating the heat release from the onset of relaxation (T onset) to the end of the relaxation spectrum (near to Tg). ΔHrel is the excess enthalpy of the glass over its relaxed state and can be used to quantify the extent of rejuvenation or aging induced by different treatments. For each set of UH conditions, three to five samples were processed and then used for DSC; in this way, the reproducibility of the results could be assessed, and a standard deviation was derived for the value of ΔHrel.

For a UH processing time of 1.0 s, the effects of static pressure are diverse [Fig. 2(b)]. In Zr55Al10Ni5Cu30 MG, ΔHrel rises at σ₀ = 3 MPa by as much as ~60% compared to the value in the as-cast glass (an increase comparable to rejuvenation by room-temperature elastostatic loading and cryogenic thermal cycling) and decreases by as much as ~27% (aging) at σ₀ = 64 MPa, ultimately tending to an enthalpy similar to that of the as-cast state. The Vit. 1 MG shows an initial sharp increase in ΔHrel followed by oscillations between states variously rejuvenated with respect to the as-cast glass. This non-monotonic trend persists even on increasing to σ₀ = 400 MPa.

![Image](image-url)
[Fig. 2(c)], and under larger pressures, $\Delta H_{rel}$ holds its value, attributable to overloading and negligible hammering effects. The $\Delta H_{rel}$ data in Figs. 2(b) and 2(c) suggest that the structural evolution induced by UH is generally non-monotonic.

The effects of UH are compared for three rods of Vit. 1 cast at different cooling rates [Fig. 2(d)]. The variation of $\Delta H_{rel}$ as a function of $r_0$ in each case is non-monotonic but correlated with the starting condition. The slowest cooled (most relaxed; $\Delta H_{rel} = 504 \text{ J mol}^{-1}$) glass shows the greatest rejuvenation on UH treatment. The fastest cooled glass ($\Delta H_{rel} = 591 \text{ J mol}^{-1}$) ultimately shows aging. The intermediate glass ($\Delta H_{rel} = 547 \text{ J mol}^{-1}$) shows moderate rejuvenation. Thus, as expected, the direction of the structural change (aging or rejuvenation) is affected not only by the conditions under which the glass is treated but also by its initial condition: a relaxed initial glass is more likely to be rejuvenated and vice versa.

We test whether non-monotonic behavior (Fig. 2) is systematic by correlating $\Delta H_{rel}$ with other properties. Prior to the DSC runs [Fig. 2(b)] on Zr$_{55}$Al$_{10}$Ni$_{5}$Cu$_{30}$ and Vit. 1 glasses, the Vickers microhardness of the disks was measured on the UH-struck surface. For this glass, higher HV is correlated with lower $\Delta H_{rel}$ [see Fig. 3(a)], as expected for a denser, more relaxed glass. Furthermore, the hardness of the Zr$_{55}$Al$_{10}$Ni$_{5}$Cu$_{30}$ MG disk treated at the static stress of 3 MPa was measured on the opposite surface (i.e., opposite to the struck surface) and along the cross section of the disk [Fig. 3(b)]. It can be observed that the hardness values on these surfaces are very close to the hardness value of the UH-struck surface (HV $\sim 480$), implying that UH has influenced the whole of the sample homogeneously.

Further correlations were explored for Vit. 1 MG. In annealed samples, UH leads to oscillations in shear modulus as a function of $r_0$ at a given hammering time [Fig. 4(a)]. When the UH process intensities, the treated samples show compressive plasticity sometimes lower, sometimes higher than that of the as-annealed glass [Figs. 4(b) and 4(c)]. Figure 4(d) shows that the higher modulus is well correlated with lower plasticity (as expected for a denser, more relaxed glass). These correlated property changes force the conclusion that there is oscillatory energy storage as a function of the extent of UH.

Even simple thermal annealing of melt-spun MG ribbons appears capable of giving oscillatory changes in properties, but these effects are complicated by possible evolving structural anisotropy (and anisotropic dimensional changes) in the samples. The broad distribution of relaxation times in MGs can lead to non-monotonic (but not oscillatory) property changes after two-stage annealing. However, the TMP of MGs has been expected to give monotonic changes.

The oscillatory response of glasses is frequently seen in viscous flow of the supercooled liquid state above $T_g$. At fixed temperature and imposed strain rate, the stress rises and shows a maximum as viscous flow starts. At higher strain rates, this stress overshoot can be followed not by monotonic decay to the steady-state flow stress but rather by stress oscillations. This effect, corresponding to oscillatory...
relaxation of the viscosity, was first seen for glassy polymers, then for MGs, and for a borosilicate glass.

The observed oscillations in MG flow stress have been modeled in terms of changes in the structural relaxation time $\tau$ in the supercooled liquid. During flow at a constant strain rate, from maximum to minimum flow stress corresponds to a decrease in $\tau$ by a factor of $\sim 5$, i.e., the relative rejuvenation that would be achieved by forming the glass on cooling times faster. In present work, the extent of the $\Delta H_{\text{fisal}}$ variation induced by UH varies widely, but it is as high as 500 J mol$^{-1}$ for Zr$_{55}$Al$_{25}$Ni$_{15}$Cu$_{10}$ MG. This is $\sim 5\%$ of the heat of melting and would correspond to the degree of rejuvenation (or aging) associated with a factor of 100 change in the cooling rate at which the glass was formed. Therefore, the amplitude of oscillations realized in a MG in the elastic regime can exceed that seen in viscous flow.

The flow behavior of materials under oscillatory excitations can be delineated by the Pipkin diagram. Figure 5 shows the Pipkin map for the flow behavior of MGs excited by various routes of oscillatory loading. The diagram illustrates strain amplitude as a function of the Deborah number ($De$). $De$ is the ratio of the relaxation time to the observation time, and for oscillatory loading conditions, $De$ can be expressed as:

$$De = \tau \omega,$$

where $\tau$ is the material relaxation time and $\omega$ is the frequency of oscillation. When $De \ll 1$, the material has a liquid-like (Newtonian) flow behavior, and when $De \gg 1$, the material has a purely elastic behavior. In the moderate values of $De$, the rheological nature of material’s flow is linear/nonlinear viscoelasticity depending on the amplitude of oscillations.

During the low amplitude oscillatory excitation of MGs such as dynamic mechanical spectroscopy (DMS), the common strain amplitudes used are quite small (typically $\sim 0.01\%$), and MGs show linear viscoelastic behavior ($De$ for La$_{60}$Ni$_{15}$Al$_{25}$ MG is between $\sim 10^{-2}$ and $\sim 10^0$). However, when a high field excitation is applied, as in large amplitude oscillatory spectroscopy (LAOS) of Pd$_{40}$Ni$_{40}$P$_{20}$ MG where $De > 1$, the viscoelastic response would be nonlinear, and the deviation from linear elasticity (Hooke’s law) becomes evident. In our UH experiments, the strain amplitudes range from $\sim 0.65\%$ to $\sim 0.9\%$ (see the supplementary material for strain considerations), and these values are quite larger than the common strain amplitudes set during DMS and LAOS excitations. Furthermore, $De$ values for UH excitation of Vit. 1 and Zr$_{55}$Al$_{25}$Ni$_{15}$Cu$_{10}$ MGs are estimated to be lower than unity. It has been proposed that MGs show serrated (nonlinear) flow behavior when $De < 1$. Therefore, UH excitation lies in the nonlinear viscoelasticity regime of the Pipkin map where nonlinear responses such as oscillatory or serrated behavior are expected.

In summary, the high-intensity processing of MGs within the elastic regime could induce both rejuvenation and aging during the same treatment in a pattern of energy storage oscillations. Consistent with the structural oscillations, mechanical properties also show oscillatory changes. The findings stress the importance of considering possible effects of energy storage oscillations in optimizing the TMP of MGs to broaden the range of available energy states and have significance for the interpretation and tuning of property changes in MGs.

See the supplementary material for the detailed description on the sample preparation, characterization procedures, and estimations of strain amplitude and Deborah number.

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