

Broadband nanoindentation of glassy polymers:

Part II. Viscoplasticity

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The relationship between hardness and flow stress in glassy polymers is examined. Materials studied include poly(methylmethacrylate), polystyrene, and polycarbonate. Properties are strongly rate dependent, so broadband nanoindentation creep (BNC) is used to measure hardness across a broad range of indentation strain rates (10^{-4} to 10 s^{-1}). Molybdenum (Mo) is also studied to serve as a “control” whose rate-dependent hardness properties have been measured previously and whose flow stress, unlike the polymers, is pressure insensitive. The BNC hardness data are converted to uniaxial flow stress using two methods based on the usual Tabor–Marsh–Johnson correlation. With both methods the resulting BNC-derived uniaxial flow stress data agree closely with literature compression uniaxial flow stress data for all materials. For the polymers, the BNC hardness data depend on initial rate of loading, indicating that the measured properties are path dependent. Path dependence is not detected in Mo.

I. INTRODUCTION

Polymers are noted for their pronounced time- and temperature-dependent mechanical properties, including yield. Rate and temperature effects in the yield stress are a consequence of the mechanisms for plastic flow, which involve thermally activated chain rearrangements and side group motions.^{1,2} Although rate and temperature effects are important in their own right, they can also be used as “fingerprints” to help gain insight into deformation mechanisms and polymer structure (e.g., Refs. 3–7). Approaches involving uniaxial compression, tension, or shear experiments are well established for bulk specimens to assess the temperature and rate dependence of yielding.

It is desired to extend studies of rate and temperature dependence of yielding to microscopic polymer systems including thin films^{8,9} and components in biological materials.^{10,11} One way to do this is to rely on indentation creep, which has been used for more than 50 years in polymers.^{12,13} Indentation creep involves pushing a hard (typically diamond), pyramid- or sphere-shaped indenter into the specimen until a predetermined load is reached, then holding the load constant. The constant load portion of the experiment is analyzed. At constant load, the indenter continues to

penetrate so that area of contact grows as a function of time. The pressure, or hardness, beneath the indenter and the rate of penetration both decrease with time giving rise to a spectrum of hardness versus strain rate. A similar experiment is impression creep in which the end of a right circular cylinder is used.¹⁴ Because the area stays constant during creep, the experiment is less ambiguous to interpret, but the load must be varied to achieve different levels of stress.

With advances in instrumentation over the past 20 years (nanoindentation), indentation creep can now readily be applied to probe rate effects in submicron volumes of material. However, there remains a lack of consensus on how best to perform and interpret indentation creep experiments in polymers. A common practice is to fit creep data to specific mathematical equations or spring and dashpot models, and then to back out material parameters from the fits.^{15–18} Some analyses assume the deformations are viscoelastic,^{19–25} whereas others account for both viscoelastic and viscoplastic deformations.^{26,27} The distinction between viscoelasticity and viscoplasticity in polymers can be gradual, and the problem of separating the two is compounded when a Berkovich indenter is used, whose characteristic strain of about 7–8% is comparable to the strain at the onset of yield. Yet, Berkovich indenters are readily available, convenient to use, and because they are self-similar they can be used to probe properties across a wide range of length scales while maintaining the average strain beneath the indenter constant.

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Our goal is to develop methods for obtaining robust and unambiguous measurements of viscoelasticity and viscoplasticity with Berkovich indenters without having to constrain the analyses by relying on constitutive models or having to make unverified assumptions of material behavior. In the accompanying article,²⁸ we evaluated viscoelastic moduli from unloading slopes of nanoindentation traces in glassy polymers. Here we assess viscoplasticity in the same materials by using the constant load creep portion of the nanoindentation trace. It turns out that the experiments and analysis require minimal assumptions about material behavior, and the connection between rate-dependent hardness and rate-dependent yield stress becomes immediately apparent when the two are compared across a wide range of strain rate.

For each instant in time during indentation creep, instantaneous Meyer hardness is defined as the average pressure beneath the indenter,

$$H = \frac{P}{A}, \quad (1)$$

where P is the applied load and A is the instantaneous projected contact area. The material strain rates beneath the indenter fall across a distribution. The median of this distribution scales in proportion to indentation strain rate, which we define as²⁹

$$\dot{\epsilon}_H = \frac{d \ln \sqrt{A}}{dt}. \quad (2)$$

In the present work, we measure the H – $\dot{\epsilon}_H$ properties of the glassy polymers poly(methylmethacrylate) (PMMA), polystyrene (PS), and polycarbonate (PC) using broadband nanoindentation creep (BNC)³⁰ to access as wide a range of strain rate as possible (10^{-4} to 10 s^{-1}). To better understand the physical significance of H , we present two methods for converting H – $\dot{\epsilon}_H$ to compression yield stress–strain rate (σ – $\dot{\epsilon}$). With both methods, the σ – $\dot{\epsilon}$ derived from BNC agrees well with literature data. This means the same viscoplastic data can be obtained from indentation creep and uniaxial experiments. To make sure we are properly analyzing creep data from the polymers, we also perform BNC experiments on molybdenum (Mo), which serves as a control because its properties have been measured before, although not over such a wide range of strain rate,^{31,32} and because its flow stress is less sensitive to hydrostatic pressure than those of the polymers.

II. EXPERIMENTAL WORK

The accompanying article²⁸ provides complete details of the experimental work. Except where noted, all experimental procedures are the same.

A. Specimens

The same PMMA, PC, and PS glassy polymers prepared in the accompanying article are also used in this

work. In addition, bulk Mo is studied. The Mo specimen was the same one studied previously by Stone and Yoder.³¹ The surface of this specimen was freshly prepared by metallographic grinding and polishing down to 0.05- μm alumina.

Before performing BNC experiments, we determined how the hardness and modulus depend on load in these materials. The motivation was to test for any depth dependence in material properties, which if present needs to be accounted for in the analysis of indentation creep.³¹ The results shown in Fig. 1 are based on areas measured using atomic force microscope (AFM) images. Mo alone shows an indentation size effect in hardness over the range

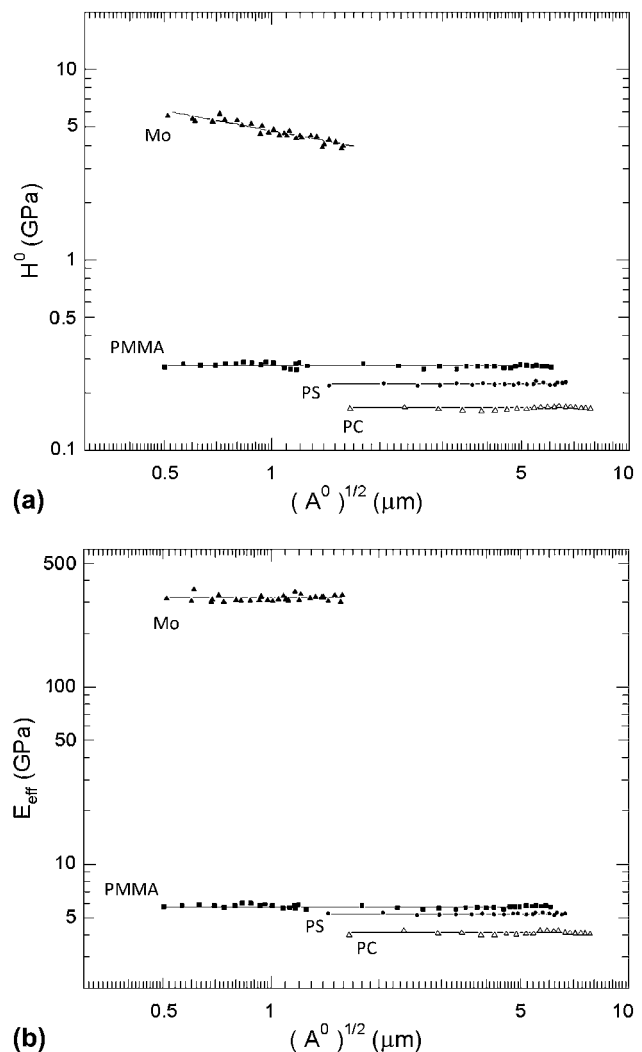


FIG. 1. Dependence of (a) Meyer hardness and (b) modulus on indent size. H^0 is the hardness immediately before unloading. Modulus (E_{eff}) was measured using methods described in the accompanying article.²⁸ The load function used to obtain these data was the same as in Fig. 1 in the accompanying article²⁸ with 0.01-s load, 10-s creep hold, and 4-s unload. For these experiments, the load, creep hold, and unload times were maintained constant, independent of maximum load, so that H^0 at different loads would correspond to the same strain rates.

of loads tested. None of the materials tested exhibit a size effect in modulus. A method for checking the area measurements (and thereby verifying the presence or absence of indentation size effect in properties) is to determine machine compliance using two different methods: Doerner–Nix,³³ which requires knowledge of indent areas, and Stone–Yoder–Sproul correlation,^{34–36} which does not. The indentation size effect is well known in large-grained metals.^{37,38} There is less consensus on the behavior of glassy polymers at shallow depths. Some researchers have reported an indentation size effect for depths less than 100 nm (e.g., Refs. 39 and 40), but this depth is shallower than any of our experiments except for the shallowest indent in PMMA.

B. Nanoindentation creep procedure

A Hysitron (Minneapolis, MN) TI 900 TriboIndenter equipped with a Berkovich probe and upgraded with a performech controller was used. Procedures concerning data acquisition rates, thermal drift, machine compliance, and corrections for the first-order filter acting on the load signal are described in the accompanying article.²⁸ Experiments were performed with open-loop control. The basic load function used is described in the accompanying article.²⁸ The only difference in the experiments described here is that we also vary the loading time, whereas in the viscoelasticity experiments loading time was maintained constant. In each material a series of indents possessing different combinations of load, creep hold, and unload times was placed. Indents began with 2-s load or “preload” to 0.46 mN, which allowed the load–depth traces from different indents to be aligned to reduce scatter in the area–depth data. Preloads were then followed by a loading ramp to 10 mN. Load times were varied between 0.01 and 100 s to identify whether loading rate affects the hardness–strain rate behavior during the subsequent creep hold. Likewise, the creep hold time at maximum load was varied between 0 and 100 s to assess how the areas of indents grow during the creep hold. Finally, unloading time was varied between 0.01 and 100 s with the purpose of measuring viscoelastic moduli, as described in the accompanying article.²⁸ The analysis of creep is not affected by unloading time provided that the area does not grow irreversibly during unloading. Indents for which this happened were discarded from the analysis.

One might reasonably ask whether adiabatic heating⁴¹ would raise the temperature at high rates of deformation and thereby alter the properties of the polymers measured using BNC. It turns out that in this regard nanoindentation experiment has advantages over conventional experiments on bulk specimens. An experiment can be considered isothermal⁴² if

$$\dot{\gamma} << \frac{2\kappa}{L^2}, \quad (3)$$

where $\dot{\gamma}$ is a shear strain rate, κ is the thermal diffusivity, and L is the characteristic length of the experiment. For the experiments reported below on polymers, the largest indents are about 10 μm across ($\approx L$). A representative value of κ is $1 \times 10^{-7} \text{ m}^2 \cdot \text{s}^{-1}$.⁴³ Indentation experiments for 10- μm indents in polymers can therefore be considered isothermal for strain rates sufficiently below 2000 s^{-1} , as was true for our experiments.

C. AFM to measure indent areas

A Quesant (Agoura Hills, CA) AFM incorporated in the TriboIndenter was used to image all residual indents. The AFM was operated in contact mode and calibrated as described in Ref. 28. ImageJ (<http://rsb.info.nih.gov/ij/>) image analysis software was used to manually measure the contact areas, A^0 , used in the analyses from 15- μm field-of-view images for the polymers and 4- μm field-of-view images for the Mo. The procedures and criteria for measuring area are described in the accompanying article.²⁸

III. EXPERIMENTAL RESULTS

A. Preliminaries

Normalized depth–time data for indents with 0.01-s load times are shown in Fig. 2 for each material tested. The depths increase smoothly with time during the creep hold. It is remarkable that in terms of normalized depth, Mo exhibits as much creep as both PS and PC. It turns out that all these materials possess very different viscoplastic behaviors, but those differences are easier to see in the area–time and hardness–indentation strain rate data than in the normalized depth–time data.

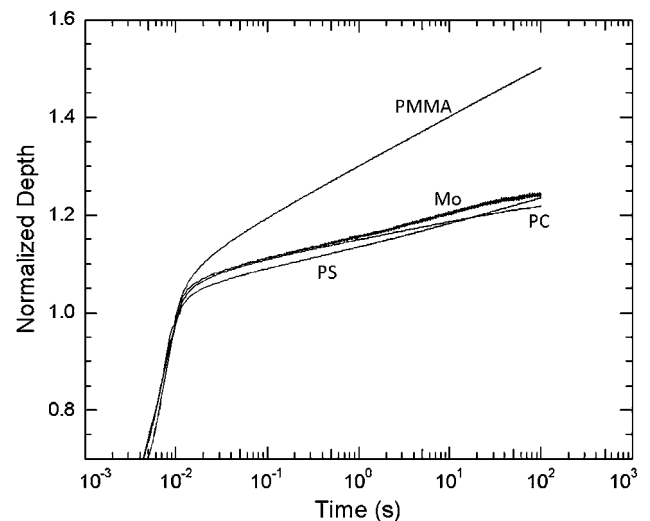


FIG. 2. Normalized depth–time traces for each material with 0.01-s load times and 100-s creep hold times. The depths are normalized so that 1 corresponds to the end of the loading ramp and beginning of the creep hold, corresponding to time = 0.01 s.

Continuous area–time curves are needed to generate instantaneous H and $\dot{\epsilon}_H$. Increase in area during the creep hold was assessed by first measuring areas from AFM images of indents with different creep hold times (Fig. 3). Normalized area–time data from experiments with 0.01-s load times are shown in Fig. 4 for each material tested. Multiple data points for a given creep hold time correspond to measurements taken from indents with different unloading times. Next, the discrete area–time measurements were fit with continuous area–time curves constructed from nanoindentation load–depth–time traces using a physically based model,³⁴ which is described in more detail in the “Appendix.” After area–time curves were constructed, Eqs. (1) and (2) were used to generate H and $\dot{\epsilon}_H$, respectively.

B. Hardness–indentation strain rate

Experimental BNC hardness–indentation strain rate (H – $\dot{\epsilon}_H$) data for PMMA, PS, PC, and Mo are shown in Figs. 5–8, respectively. Also included are compression flow stress–strain rate (σ – $\dot{\epsilon}$) data taken from the literature^{42,44} and σ – $\dot{\epsilon}$ converted from H – $\dot{\epsilon}_H$ using method 1 described below. For now we will ignore the σ – $\dot{\epsilon}$ data and are only concerned with the H – $\dot{\epsilon}_H$ data in Figs. 5–8. BNC H – $\dot{\epsilon}_H$ data with 0.01-s load times span strain rates between approximately 10^{-4} and 10 s^{-1} , whereas experiments with longer load times span narrower ranges of strain rate because the creep portions of those experiments begin at lower strain rates. Also included in the polymer plots are H estimated from 0-s creep hold indents. These indents had 0.01-s load and unload times. Previous authors⁴⁶ noted that the strain rate during a loading ramp is approximately $\dot{P}/2P$, where $\dot{P}$ is the rate of loading. Loading 10 mN in 0.01 s corresponds to a strain rate of approximately 50 s^{-1} immediately after the 0.01-s loading ramp, so these data are placed at this strain rate. The areas of indents with 0-s hold times likely grew irreversibly during unloading, so these values represent an approximate lower bound for H at this strain rate.

It can be seen from Figs. 5–7 that for the polymers, H – $\dot{\epsilon}_H$ behavior depends on initial rate of loading. At fixed $\dot{\epsilon}_H$, the H is higher for slower loading rates. The fact that the properties depend on how they are measured means that they are path dependent. Path-dependent behavior is not observed in Mo (Fig. 8) nor in BNC of metallic glasses³⁰ also tested at room temperature. We surmise that for the polymers, rate- and time-dependent structural changes take place during the loading ramp and that these changes affect the hardness during the subsequent creep hold. The observed behavior is likely related to the yield drop seen in compression tests of amorphous polymers.

The hardness in Mo exhibits an indentation size effect (ISE) [Fig. 1(a)]. Therefore, changes in H during indentation creep derive from changes in both strain rate and depth³¹ following

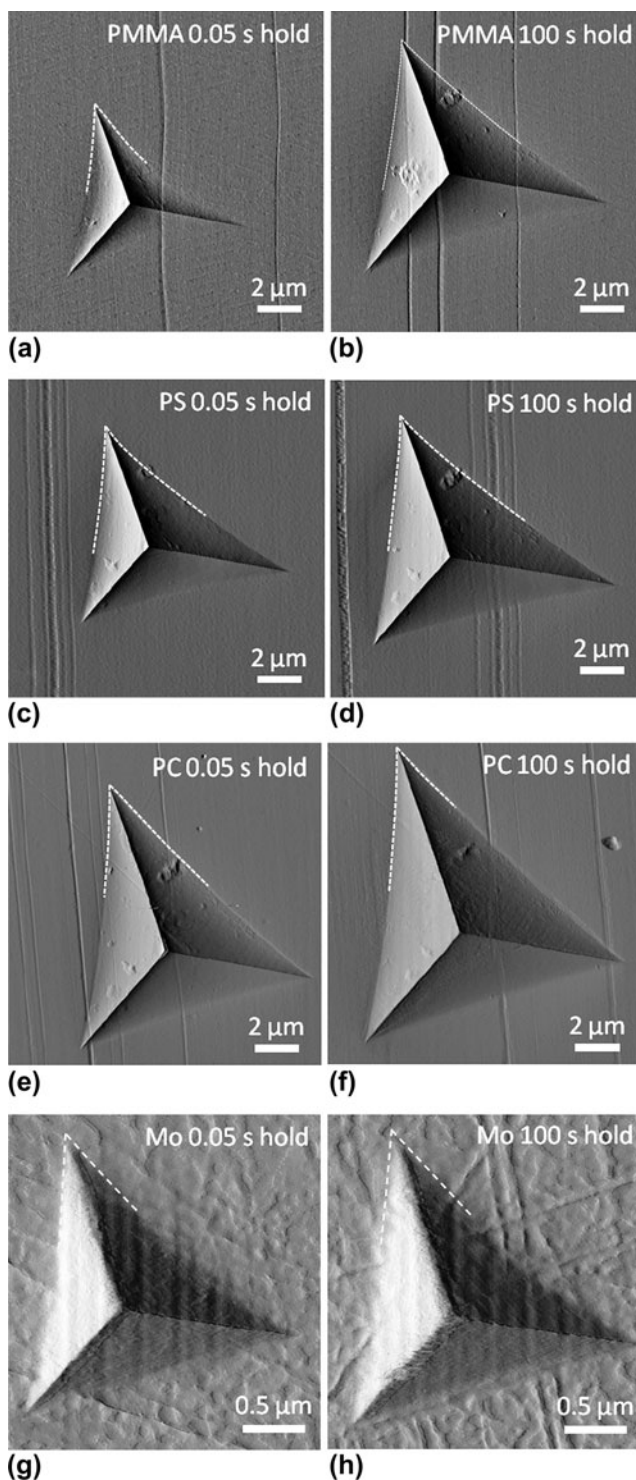


FIG. 3. Atomic force microscopy (AFM) images of 10-mN indents with 0.05- and 100-s creep hold times in (a and b) poly(methylmethacrylate) (PMMA); (c and d) polystyrene (PS); (e and f) polycarbonate (PC); and (g and h) molybdenum (Mo). All indents had 0.01-s load and unload times. Dashed lines highlight the contact edges used to assess A^0 . The identification of the contact edges for the polymers was described in detail in the accompanying article.²⁸

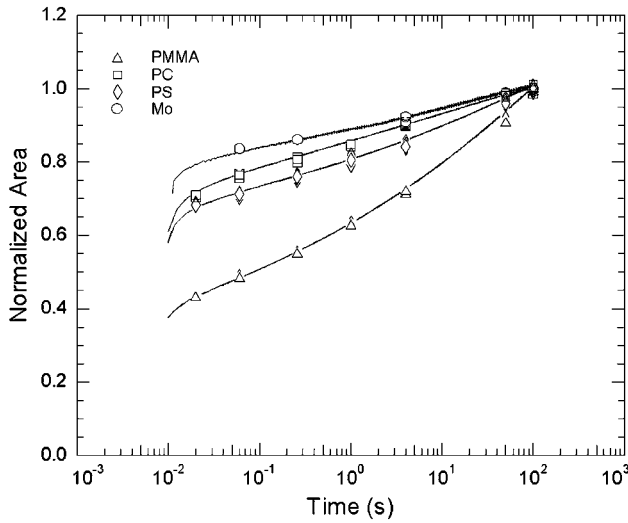


FIG. 4. Normalized area–time data from 100-s creep hold time for each material tested. The areas are normalized so that 1 corresponds to the area at the end of the 100-s creep hold time. The beginning of creep hold of the experiment is 0.01 s. The discrete points were measured from AFM images. The continuous curves were generated with the physically based model³⁴ described in the “Appendix.”

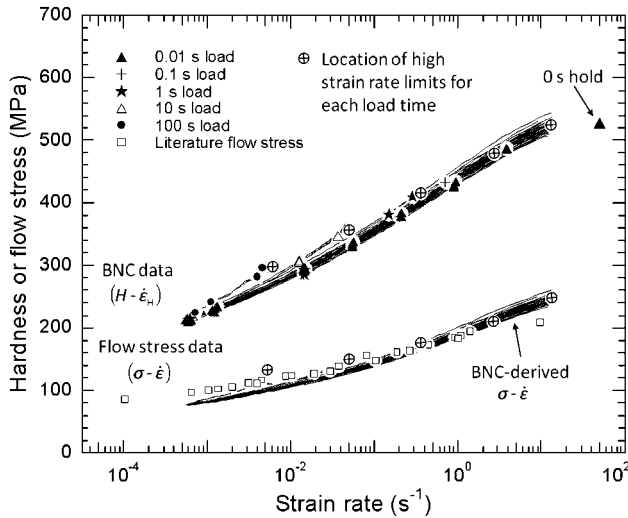


FIG. 5. PMMA broadband nanoindentation creep (BNC) data ($H-\dot{\epsilon}_H$) collected for load times ranging from 0.01 to 100 s. The high strain rate limits for each load time are represented by circles with cross hairs. Also included are two sets of flow stress data ($\sigma-\dot{\epsilon}$): (i) literature $\sigma-\dot{\epsilon}$ from uniaxial compression tests (Lee and Swallowe⁴² and Richeton et al.⁴⁴) and (ii) BNC-derived $\sigma-\dot{\epsilon}$ calculated using the correlation in Fig. 9 following method 1 described in the “Discussion.”

$$\Delta \ln H = m_H \Delta \ln \dot{\epsilon}_H + n \Delta \ln h_p \quad , \quad (4)$$

where m_H is the strain rate sensitivity of the hardness ($\partial \ln H / \partial \ln \dot{\epsilon}_H|_{\text{fixed depth}}$) and n measures the strength of the size effect. We measured $n = -0.35$ from the data in Fig. 1(a). Because of our interest in changes in H that derive solely from changes in $\dot{\epsilon}_H$, we subtract the second term on the right-hand side of Eq. (4) from experimental

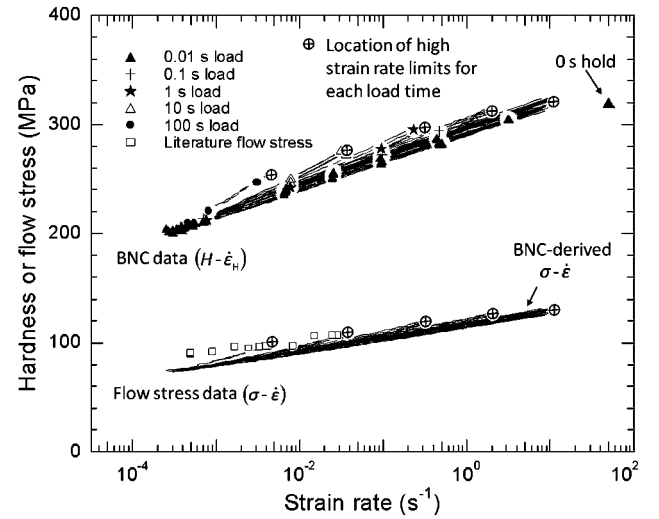


FIG. 6. PS BNC data ($H-\dot{\epsilon}_H$) collected at load times ranging from 0.01 to 100 s. The high strain rate limits for each load time are represented by circles with cross hairs. Also included are two sets of flow stress data ($\sigma-\dot{\epsilon}$): (i) literature $\sigma-\dot{\epsilon}$ from uniaxial compression tests (Lee and Swallowe⁴²) and (ii) BNC-derived $\sigma-\dot{\epsilon}$ calculated using the correlation in Fig. 9 following method 1 described in the “Discussion.”

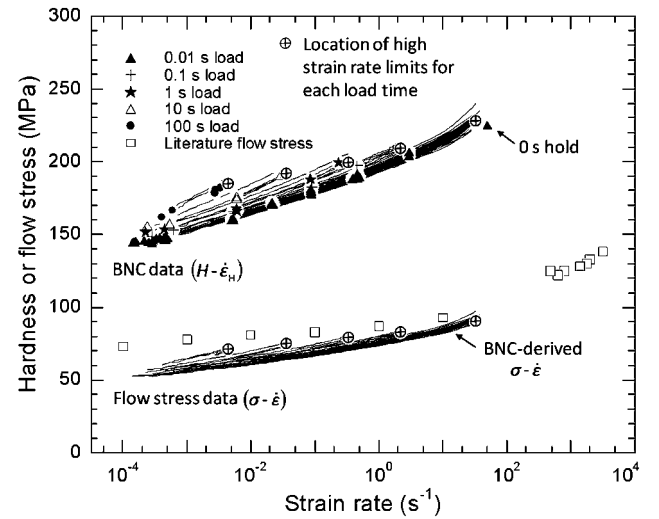


FIG. 7. PC BNC data ($H-\dot{\epsilon}_H$) collected at load times ranging from 0.01 to 100 s. The high strain rate limits for each load time are represented by circles with cross hairs. Also included are two sets of flow stress data ($\sigma-\dot{\epsilon}$): (i) literature $\sigma-\dot{\epsilon}$ from uniaxial compression tests (Richeton et al.⁴⁴) and (ii) BNC-derived $\sigma-\dot{\epsilon}$ calculated using the correlation in Fig. 9 following method 1 described in the “Discussion.”

data to give H as a function of $\dot{\epsilon}_H$ at fixed depth. Hardness– $\dot{\epsilon}_H$ data with and without the correction for the indentation size effect are shown in Fig. 8.

IV. DISCUSSION

A. Conversion of hardness to flow stress

It is of interest to map BNC $H-\dot{\epsilon}_H$ data onto literature $\sigma-\dot{\epsilon}$ data. Two methods produce satisfactory results. Both

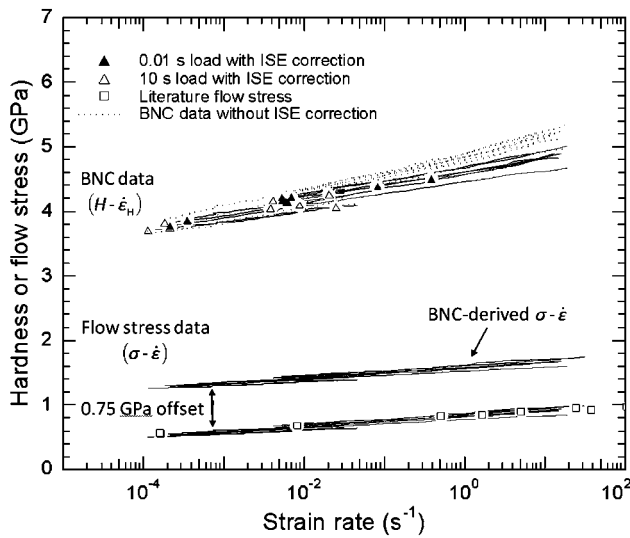


FIG. 8. Mo BNC data ($H-\dot{\epsilon}_H$) collected at load times of 0.01 and 10 s. In contrast to the polymers, Mo does not exhibit path dependence in H , but it does exhibit an indentation size effect (ISE) in H . Hardness- $\dot{\epsilon}_H$ curves with the ISE correction [Eq. (4)] are plotted as solid lines, whereas the $H-\dot{\epsilon}_H$ curves before ISE correction are plotted as dotted lines. Also included are three sets of flow stress data ($\sigma-\dot{\epsilon}$): (i) literature $\sigma-\dot{\epsilon}$ from uniaxial compression tests (Briggs and Campbell⁴⁵), (ii) BNC-derived $\sigma-\dot{\epsilon}$ calculated using the correlation in Fig. 9 following method 1 described in the “Discussion,” and (iii) BNC-derived $\sigma-\dot{\epsilon}$ offset by 0.75 GPa to account for differences in work hardening.

rely on the Tabor–Marsh–Johnson correlation^{47–51} in which $H = k\sigma$ and k is a function of indenter angle (α in Fig. 9) and E^* / σ , where E^* is the Hertzian contact modulus defined in Eq. (A2) of the “Appendix.” The difference between the two methods resides in how they convert strain rate. The first method is simpler: it assumes that $\dot{\epsilon}_H = k'\dot{\epsilon}$, where $\dot{\epsilon}$ is the strain rate for a uniaxial compression test and k' is a constant, independent of hardness (we use $k' = 1$). This method was proposed by Kermouche et al.^{52,53} and Elm Mustafa et al.²⁹ The second method, preferred by the latter authors because it does a better job of fitting finite element simulations, relaxes the assumption that k' is constant. The two methods are virtually identical for low H / E^* ratio where $k' = \text{constant}$ is a good approximation, and their difference increases as H / E^* increases. For the second method, $\dot{\epsilon}$ can be generated by first taking m_H and converting it to strain rate sensitivity of the flow stress ($m_\sigma = \partial \ln \sigma / \partial \ln \dot{\epsilon}$) using the ansatz

$$\frac{m_H}{m_\sigma} \approx \sqrt{1 - (x/0.18)^2} \times \left[0.91 - 0.057 \arctan\left(\frac{\log(x/24.8)}{0.2}\right) \right], \quad (5)$$

where $x = H / E^*$. The parameters used in Eq. (5) differ from the earlier version^{29,30} based on finite element analysis, although the difference does not drastically

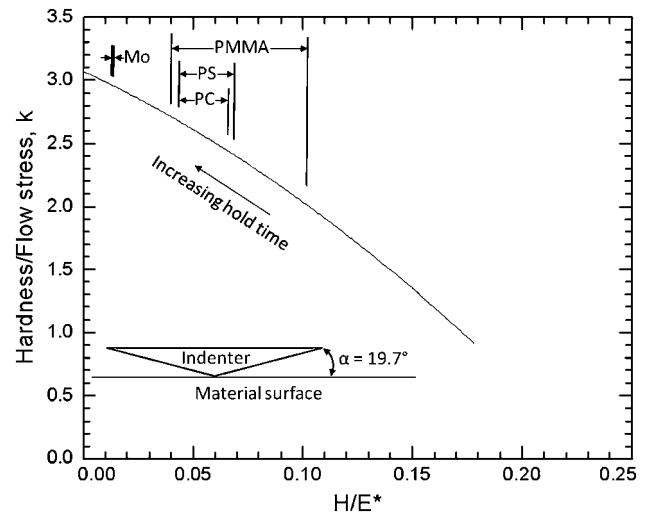


FIG. 9. The value of k as a function of H/E^* for a Berkovich probe (cone equivalent $\alpha = 19.7^\circ$) based on analysis of the Tabor–Marsh–Johnson correlation.⁵¹

change the results. The new parameters have been motivated by discussions with A.A. Elm Mustafa (personal communication), who is modeling indentation creep for a variety of indenter geometries including Berkovich. In the earlier version, 0.21 was used as normalization for x inside the square root sign rather than 0.18, and 28.2 was used inside the arctangent function rather than 24.8 applied now. The value of $x = 0.21$ corresponds to the point at which $m_H / m_\sigma \rightarrow 0$ and was obtained empirically based on finite element analysis. Not surprisingly, it turns out that $x = 0.21$ also corresponds to the fully elastic limit based on Sneddon’s theory,⁵⁴ which gives $H / E^* = 0.202$ indentation of an elastic solid with a 22° cone. The new expression therefore takes into account the difference between the cone angle (22°) for the simulations and equivalent cone angle (19.7°) for a Berkovich indenter. It assumes that both of the normalizing factors for x scale in proportion to $\tan \alpha$. Strain rate is obtained by integrating inverse strain rate sensitivity,

$$\ln \dot{\epsilon} = \int_{\ln \sigma_{\min}}^{\ln \sigma} \frac{1}{m_\sigma(s)} d \ln s + \ln \dot{\epsilon}_{\min}, \quad (6)$$

in which $\ln \dot{\epsilon}_{\min}$ is an integration constant with $\dot{\epsilon}_{\min} = \dot{\epsilon}_H / k'$ evaluated at the lowest hardness in the BNC experiment ($H_{\min}$), corresponding to the lowest stress, $\sigma_{\min}$. From Ref. 29, we estimate $k'(\sigma_{\min}) = 2$ for PMMA, 3.3 for PS, and 3.5 for PC.

The ratio k is shown in Fig. 9 for the Berkovich geometry and as a function of H / E^* .³⁰ To convert H to σ , H / E^* must be estimated at each point in the BNC data. The hardness is known, but for the polymers the moduli lack uniqueness because of viscoelasticity. We have yet to

perform a systematic investigation of how best to deal with the viscoelastic modulus in this analysis. Nevertheless, the results are not too sensitive to the value of modulus chosen, and for the conversion here, we arbitrarily choose a value of E^* that corresponds to the average measured from experiments with 1-s unloading time.²⁸ The ranges of k values utilized for converting different BNC data are shown in Fig. 9.

The BNC-derived σ - $\dot{\epsilon}$ curves constructed using method 1 are compared with literature σ - $\dot{\epsilon}$ data in Figs. 5–8. For the polymers, the high strain rate limits of the BNC-derived σ - $\dot{\epsilon}$ curves agree quite well with the literature data in Figs. 5–7. This is probably because the high strain rate data from BNC correspond to the beginning of the creep hold and to the behavior of the freshly deformed material, analogous to the onset of yielding in uniaxial compression, whereas the remainder of the creep hold, which agrees less well with the literature yield stress data, assesses properties affected by prior deformation, similar to the lower yield stress in a uniaxial compression experiment. The second method of converting BNC data tends to elongate the stress-strain rate data in the direction of strain rate (Fig. 10). The clearest difference in outcome between the two methods is for PC, where method 2 more accurately captures the upward curvature of the σ - $\dot{\epsilon}$ data at high strain rates.

The same method for converting BNC data for polymers can be used for Mo. In this case $k \approx 3$ (Fig. 9). In contrast to the BNC-derived σ - $\dot{\epsilon}$ data for the polymers, the σ - $\dot{\epsilon}$ data for Mo fall at higher levels of stress than the literature σ - $\dot{\epsilon}$ data. This result is not surprising. Mo both work hardens and exhibits an indentation size effect, so we do not necessarily expect BNC-derived σ - $\dot{\epsilon}$ to agree with literature σ - $\dot{\epsilon}$, which correspond to a different level of work hardening. Instead, the derivative $d\sigma/d\log \dot{\epsilon}$, which is

independent of work hardening,^{31,45} agrees. When the nanoindentation-derived σ - $\dot{\epsilon}$ data are offset downward in Fig. 8 by 0.75 GPa, they overlap the literature data.

B. Role of hydrostatic pressure in conversion of hardness to flow stress

Hydrostatic pressure is known to affect yielding in polymers,^{55–57} which, for instance, is often modeled using Drucker–Prager or Mohr–Coulomb-type laws. A number of empirical and theoretical studies suggest that for materials that are sensitive to hydrostatic pressure, the ratio H/σ is increased.^{58–62} In our analysis we have not directly accounted for hydrostatic stress but have relied solely on the empirical relationship shown in Fig. 9. However, much of the experimental data that went into constructing this relationship came from polymers like PMMA and PS.⁴⁹ In this case the pressure dependence of yielding in polymers is already implicitly included in the correlation.

C. Comparison of current PMMA results to previous work

We had previously tried the same comparison between BNC-derived σ - $\dot{\epsilon}$ curves with literature σ - $\dot{\epsilon}$ data for PMMA.³⁰ The agreement was rather poor and was attributed to effects of hydrostatic pressure. However, the older measurements were in error. As described in the accompanying article,²⁸ we have reassessed the way we measure the areas of PMMA indents and have found we had previously underestimated the areas of indents with fast loading rates and short creep hold times. That error has been corrected here.

V. CONCLUDING REMARKS

BNC is used to generate hardness-indentation strain rate (H - $\dot{\epsilon}_H$) data from amorphous polymers PMMA, PC, and PS and from the body-centered cubic metal Mo. A key part of the analysis is the ability to measure and accurately model the evolution of areas during the creep hold. Two slightly different methods were also used to convert H - $\dot{\epsilon}_H$ data into BNC-derived σ - $\dot{\epsilon}$ data. We found the following:

(1) BNC H - $\dot{\epsilon}_H$ curves for PMMA, PS, and PC exhibit path dependence. The H - $\dot{\epsilon}_H$ curves depend on the rate of loading preceding the creep hold.

(2) In contrast, BNC H - $\dot{\epsilon}_H$ curves for Mo do not exhibit path dependence. For Mo, the H - $\dot{\epsilon}_H$ curves are independent of loading rate.

(3) The BNC H - $\dot{\epsilon}_H$ data were converted to σ - $\dot{\epsilon}$ data using two methods. Both methods rely on the Tabor–Marsh–Johnson correlation to convert H to σ . The first method assumes the $\dot{\epsilon}_H$ is strictly proportional to $\dot{\epsilon}$, whereas with the second method the strain rate sensitivity of the hardness is converted to that of the flow stress, then the

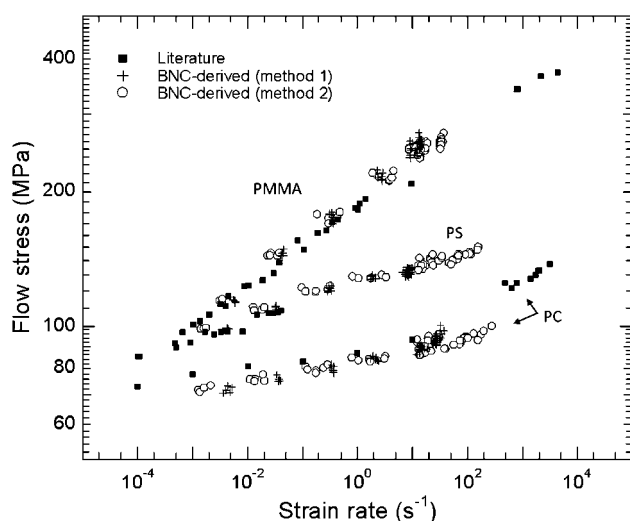


FIG. 10. Comparison between BNC-derived σ - $\dot{\epsilon}$ and literature uniaxial σ - $\dot{\epsilon}$ data for the polymers. The BNC-derived σ - $\dot{\epsilon}$ data calculated using both methods are included.

corresponding strain rates are obtained by integration. The two methods generate similar BNC-derived σ - $\dot{\epsilon}$ data, but method 2 better replicates the behavior of the literature data for PC at high strain rates.

(4) For the polymers, the high strain rate limits of the BNC-derived σ - $\dot{\epsilon}$ curves for each loading rate agree best with the literature σ - $\dot{\epsilon}$ data. This means that it is necessary to perform experiments with different loading rates to generate σ - $\dot{\epsilon}$ data from BNC data that will compare well with literature uniaxial σ - $\dot{\epsilon}$ data. For Mo, the BNC-derived σ - $\dot{\epsilon}$ data did not depend on loading rate and agree across the entire spectrum of strain rate. In this case, the σ - $\dot{\epsilon}$ data can (at least in principle) be generated in a single BNC experiment.

(5) In generating the BNC H - $\dot{\epsilon}_H$ curves, no assumptions are made about constitutive laws that govern the materials. It is nevertheless unmistakable that BNC-derived σ - $\dot{\epsilon}$ curves reflect the viscoplastic properties of the materials.

(6) The analyses to obtain BNC-derived σ - $\dot{\epsilon}$ curves rely on the ratio of H / E^* . For the polymers, E^* is not uniquely defined, so questions remain concerning how best to convert H to σ . Clearly, the overall approach is valid, but further details remain to be worked out.

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REFERENCES

1. R.F. Boyer: Dependence of mechanical properties on molecular motion in polymers. *Polym. Eng. Sci.* **8**(3), 161 (1968).
2. A.S. Argon: A theory for the low-temperature plastic deformation of glassy polymers. *Philos. Mag.* **28**(4), 839 (1973).
3. A.E. Mayr, W.D. Cook, and G.H. Edward: Yielding behaviour in model epoxy thermosets—I. Effect of strain rate and composition. *Polymer* **39**, 3719 (1998).
4. D. Rana, V. Sauvant, and J.L. Halary: Molecular analysis of yielding in pure and antiplasticized epoxy-amine thermosets. *J. Mater. Sci.* **37**(24), 5267 (2002).
5. L. Lin and A.S. Argon: Rate mechanism of plasticity in the crystalline component of semicrystalline nylon 6. *Macromolecules* **27**(23), 6903 (1994).
6. N.W.J. Brooks, R.A. Duckett, and I.M. Ward: Temperature and strain-rate dependence of yield stress of polyethylene. *J. Polym. Sci., Part B: Polym. Phys.* **36**(12), 2177 (1998).
7. T. Kazmierczak, A. Galeski, and A.S. Argon: Plastic deformation of polyethylene crystals as a function of crystal thickness and compression rate. *Polymer* **46**(21), 8926 (2005).
8. A. Strojny, X. Xia, A. Tsou, and W.W. Gerberich: Techniques and considerations for nanoindentation measurements of polymer thin film constitutive properties. *J. Adhes. Sci. Technol.* **12**, 1299 (1998).
9. O.L. Warren and T.J. Wyrobek: Nanomechanical property screening of combinatorial thin-film libraries by nanoindentation. *Meas. Sci. Technol.* **16**(1), 100 (2005).
10. D.M. Ebenstein and L.A. Pruitt: Nanoindentation of biological materials. *Nano Today* **1**(3), 26 (2006).
11. J.E. Jakes, C.R. Frihart, and D.S. Stone: Creep properties of micron-size domains in ethylene glycol modified wood across 4½ decades in strain rate, in *Mechanics of Biological and Biomedical Materials*, edited by K. Katti, C. Hellmich, U.G.K. Wegst, and R. Narayan (Mater. Res. Soc. Symp. Proc. **1132**, Warrendale, PA, 2009), 1132-Z07-21.
12. W. Rostoker and J.O. Galante: Indentation creep of polymers for human joint applications. *J. Biomed. Mater. Res.* **13**(5), 825 (1979).
13. M. Takahashi, M.C. Shen, R.B. Taylor, and A.V. Tobolsky: Master curves for some amorphous polymers. *J. Appl. Polym. Sci.* **8**(4), 1548 (1964).
14. S.N.G. Chu and J.C.M. Li: Impression creep; a new creep test. *J. Mater. Sci.* **12**(11), 2200 (1977).
15. J. Chen, G.A. Bell, D. Hanshan, J.F. Smith, and B.D. Beake: A study of low temperature mechanical properties and creep behaviour of polypropylene using a new sub-ambient temperature nanoindentation test platform. *J. Phys. D: Appl. Phys.* **43**(42), 425404 (2010).
16. M.L. Oyen: Sensitivity of polymer nanoindentation creep measurements to experimental variables. *Acta Mater.* **55**(11), 3633 (2007).
17. A.C. Fischer-Cripps: A simple phenomenological approach to nanoindentation creep. *Mater. Sci. Eng., A* **385**(1–2), 74 (2004).
18. H. Li and A.H.W. Ngan: Size effects of nanoindentation creep. *J. Mater. Res.* **19**(2), 513 (2004).
19. G. Constantinides, C.A. Tweedie, D.M. Holbrook, P. Barragan, J.F. Smith, and K.J. Van Vliet: Quantifying deformation and energy dissipation of polymeric surfaces under localized impact. *Mater. Sci. Eng., A* **489**, 403 (2008).
20. A. Jager and R. Lackner: Identification of viscoelastic model parameters by means of cyclic nanoindentation testing. *Int. J. Mater. Res.* **99**, 829 (2008).
21. H. Lu, B. Wang, J. Ma, G. Huang, and H. Viswanathan: Measurement of creep compliance of solid polymers by nanoindentation. *Mech. Time-Depend. Mater.* **7**, 189 (2003).
22. M.L. Oyen: Spherical indentation creep following ramp loading. *J. Mater. Res.* **20**(8), 2094 (2005).
23. M.L. Oyen: Relating viscoelastic nanoindentation creep and load relaxation experiments. *Int. J. Mater. Res.* **99**, 823 (2008).
24. C.A. Tweedie and K.J. Van Vliet: Contact creep compliance of viscoelastic materials via nanoindentation. *J. Mater. Res.* **21**(6), 1576 (2006).
25. M.R. Vanlandingham, N.K. Chang, P.L. Drzal, C.C. White, and S.H. Chang: Viscoelastic characterization of polymers using instrumented indentation. I. Quasi-static testing. *J. Polym. Sci., Part B: Polym. Phys.* **43**, 1794 (2005).
26. B. Beake: Modelling indentation creep of polymers: A phenomenological approach. *J. Phys. D: Appl. Phys.* **39**, 4478 (2006).
27. C.Y. Zhang, Y.W. Zhang, K.Y. Zeng, and L. Shen: Characterization of mechanical properties of polymers by nanoindentation tests. *Philos. Mag.* **86**, 4487 (2006).
28. J.E. Jakes, R.S. Lakes, and D.S. Stone: Broadband nanoindentation of glassy polymers: Part I. Viscoelasticity. *J. Mater. Res.* **27**(2), 463 (2011).
29. A.A. Elmestafa, S. Kose, and D.S. Stone: The strain-rate sensitivity of the hardness in indentation creep. *J. Mater. Res.* **22**(4), 926 (2007).
30. J.B. Puthoff, J.E. Jakes, H. Cao, and D.S. Stone: Investigation of thermally activated deformation in amorphous PMMA and Zr-Cu-Al bulk metallic glasses with broadband nanoindentation creep. *J. Mater. Res.* **24**(3), 1279 (2009).
31. D.S. Stone and K.B. Yoder: Division of the hardness of molybdenum into rate-dependent and rate-independent components. *J. Mater. Res.* **9**(10), 2524 (1994).
32. K.B. Yoder, A.A. Elmestafa, J.C. Lin, R.A. Hoffman, and D.S. Stone: Activation analysis of deformation in evaporated molybdenum thin films. *J. Phys. D: Appl. Phys.* **36**, 884 (2003).

33. M.F. Doerner and W.D. Nix: A method for interpreting the data from depth-sensing indentation instruments. *J. Mater. Res.* **1**(4), 601 (1986).
34. D.S. Stone, J.E. Jakes, J.B. Puthoff, and A.A. Elmustafa: Analysis of indentation creep. *J. Mater. Res.* **25**(4), 611 (2010).
35. J.E. Jakes, C.R. Frihart, J.F. Beecher, R.J. Moon, P.J. Resto, Z.H. Melgarejo, O.M. Surez, H. Baumgart, A.A. Elmustafa, and D.S. Stone: Nanoindentation near the edge. *J. Mater. Res.* **24**(3), 1016 (2009).
36. J.E. Jakes, C.R. Frihart, J.F. Beecher, R.J. Moon, and D.S. Stone: Experimental method to account for structural compliance in nanoindentation measurements. *J. Mater. Res.* **23**(4), 1113 (2008).
37. Q. Ma and D.R. Clarke: Size dependent hardness of silver single crystals. *J. Mater. Res.* **10**(4), 853 (1995).
38. W.D. Nix and H. Gao: Indentation size effects in crystalline materials: A law for strain gradient plasticity. *J. Mech. Phys. Solids* **46**(3), 411 (1998).
39. C.A. Tweedie, G. Constantinides, K.E. Lehman, D.J. Brill, G.S. Blackman, and K.J. Van Vliet: Enhanced stiffness of amorphous polymer surfaces under confinement of localized contact loads. *Adv. Mater.* **19**(18), 2540 (2007).
40. E. Amitay-Sadovsky, B. Ward, G.A. Somorjai, and K. Komvopoulos: Nanomechanical properties and morphology of thick polyurethane films under contact pressure and stretching. *J. Appl. Phys.* **91**, 375 (2002).
41. A.C. Fischer-Cripps: Significance of a local temperature rise in nanoindentation testing. *J. Mater. Sci.* **39**(18), 5849 (2004).
42. G. Swallowe and S. Lee: Quasi-static and dynamic compressive behaviour of poly(methyl methacrylate) and polystyrene at temperatures from 293 K to 363 K. *J. Mater. Sci.* **41**(19), 6280 (2006).
43. N.A. Fleck, W.J. Stronge, and J.H. Liu: High strain-rate shear response of polycarbonate and polymethyl methacrylate. *Proc. R. Soc. London, Ser. A* **429**, 459 (1990).
44. J. Richeton, S. Ahzi, K.S. Vecchio, F.C. Jiang, and R.R. Adharapurapu: Influence of temperature and strain rate on the mechanical behavior of three amorphous polymers: Characterization and modeling of the compressive yield stress. *Int. J. Solids Struct.* **43**(7–8), 2318 (2006).
45. T.L. Briggs and J.D. Campbell: The effect of strain rate and temperature on the yield and flow of polycrystalline niobium and molybdenum. *Acta Metall.* **20**(5), 711 (1972).
46. B.N. Lucas and W.C. Oliver: Indentation power-law creep of high-purity indium. *Metall. Mater. Trans. A* **30**(3), 601 (1999).
47. D. Tabor: A simple theory of static and dynamic hardness. *Proc. R. Soc. London, Ser. A* **192**, 247 (1948).
48. D. Tabor: The hardness of solids. *Rev. Phys. Technol.* **1**(3), 145 (1970).
49. D.M. Marsh: Plastic flow in glass. *Proc. R. Soc. London, Ser. A* **279** (1378), 420 (1964).
50. K.L. Johnson: Correlation of indentation experiments. *J. Mech. Phys. Solids* **18**(2), 115 (1970).
51. K.L. Johnson: *Contact Mechanics* (Cambridge University Press, Cambridge, United Kingdom, 1985), p. 452.
52. G. Kermouche, J.L. Loubet, and J.M. Bergheau: Cone indentation of time-dependent materials: The effects of the indentation strain rate. *Mech. Mater.* **39**, 24 (2007).
53. G. Kermouche, J.L. Loubet, and J.M. Bergheau: A new index to estimate the strain rate sensitivity of glassy polymers using conical/pyramidal indentation. *Philos. Mag.* **86**, 5667 (2006).
54. I.N. Sneddon: Relation between load and penetration in axisymmetric Boussinesq problem for punch of arbitrary profile. *Int. J. Eng. Sci.* **3**, 47 (1965).
55. L.A. Davis and C.A. Pampillo: Deformation of polyethylene at high pressure. *J. Appl. Phys.* **42**, 4659 (1971).
56. L.A. Davis and C.A. Pampillo: Kinetics of deformation of PTFE at high pressure. *J. Appl. Phys.* **43**, 4285 (1972).
57. K.D. Pae, J.A. Sauer, and A.A. Silano: Shear deformation under hydrostatic pressure of polytetrafluoroethylene and polycarbonate, in *Proceedings of the Sixth International Conference on High-Pressure Science and Technology*, Boulder, CO, edited by K.D. Timmerhaus and M.S. Barber (1979), pp. 512–518.
58. I.W. Chen: Implications of transformation plasticity in ZrO₂-containing ceramics: II. Elastic-plastic indentation. *J. Am. Ceram. Soc.* **69**, 189 (1986).
59. P. Bardia and R. Narasimhan: Characterisation of pressure-sensitive yielding in polymers. *Strain* **42**, 187 (2006).
60. V. Keryvin: Indentation as a probe for pressure sensitivity of metallic glasses. *J. Phys. Condens. Matter* **20**(11), 114119 (2008).
61. M-Q. Le and S-S. Kim: Element analysis of instrumented sharp indentations into pressure-sensitive materials. *J. Mater. Sci. Technol.* **23**(2), 277 (2007).
62. R. Narasimhan: Analysis of indentation of pressure sensitive plastic solids using the expanding cavity model. *Mech. Mater.* **36**(7), 633 (2004).

APPENDIX

We obtain the continuous area–time curves in Fig. 4 by fitting continuous curves to the discrete area–time measurements also shown in the figure. Rather than relying on arbitrary functions like polynomials, power laws, or decaying exponentials to generate continuous area–time curves, we employ another approach that utilizes the nanoindentation load–depth–time traces. The method does a better job of capturing the trend of the data over a wide range of strain rate and has been validated and improved using finite element analysis.³⁴ With this method it is assumed that $A^{1/2} \propto h_p^{\zeta_p}$ at constant load. The power law exponent ζ_p is used as a fitting parameter to fit the continuous area–time curve to the discrete experimental data. During the creep hold, h_p is estimated using

$$h_p \cong h_p^0 + \frac{h_t - h_p^0 - \frac{P}{E_{\text{eff}}(A^0)^{1/2}} \left(1 - (1 - \zeta_p) \frac{P - P^0}{2P^0} \right)}{1 - \frac{P \zeta_p}{h_p^0 E_{\text{eff}}(A^0)^{1/2}}}, \quad (\text{A1})$$

where P is the instantaneous load (ideally, $P = P^0$ although in a real experiment P varies a small amount). The superscripts “0” refer to the values of the parameters at the end of the creep hold and $h_p^0 = h_t^0 - P^0(A^0)^{1/2}/E_{\text{eff}}$. E_{eff} is given by

$$\frac{1}{E_{\text{eff}}} = \frac{1}{\beta} \left(\frac{1 - \nu_s^2}{E_s} + \frac{1 - \nu_d^2}{E_d} \right) \equiv \frac{1}{\beta E^*}, \quad (\text{A2})$$

where E_s and E_d are Young's moduli and ν_s and ν_d are Poisson's ratios of specimen and indenter, respectively, and $\beta \approx 1.23$.³⁶ E^* is the usual modulus encountered in Hertz contact problems. Equation (A1) is obtained from a theoretical analysis. However, for polymers E_{eff} lacks uniqueness because of viscoelasticity, so the polymers do not strictly conform to the theory. In practice, this discrepancy does not present a problem because ζ_p is calculated using a representative value of E_{eff} and in the end the continuous area-time curve is insensitive to the choice of E_{eff} . Figure A1 shows $\ln(A^0)^{1/2}$ versus $\ln(h_p^0)$ from experiments with different creep hold times at fixed load. The slopes of the curves correspond to ζ_p . Rather than forming straight lines, the data must be approximated as parabolas,

$$\ln(A^0)^{1/2} = X_0 + X_1 \ln(h_p^0) + X_2 (\ln(h_p^0))^2, \quad (\text{A3})$$

in which case $\zeta_p = X_1 + 2X_2 \ln(h_p^0)$. Previously³⁴ we had not been able to reliably detect that ζ_p varies with depth. However, our measurement methods are now better, and the finite element analyses neglected viscoelastic deformation, so it should come as no surprise that ζ_p is not a constant for the polymers.

Because ζ_p depends on h_p , it is not straightforward to calculate h_p using Eq. (A1). Therefore, to account for variation in ζ_p in Eq. (A1) we divide the raw depth-load-time data into 100 equal segments on a log(time) scale so

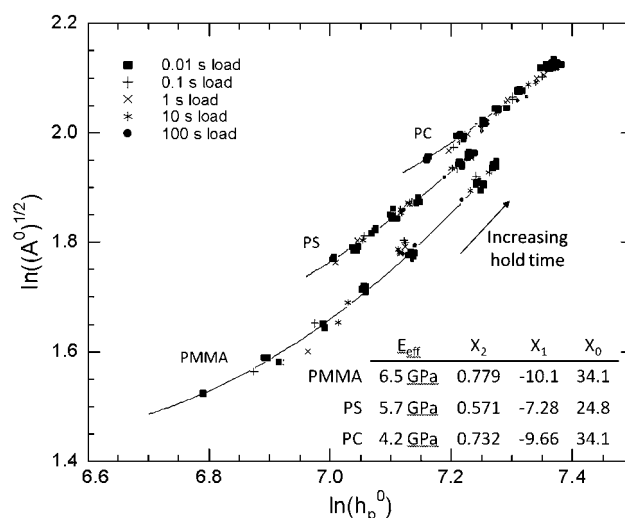


FIG. A1 Data used to calculate ζ_p for PMMA, PS, and PC. For each material, data include indents with creep hold times that vary from 0.01 to 100 s and load times that vary from 0.01 to 100 s. The 0.01-s load time data were fit to Eq. (A3), and these fits were differentiated to calculate ζ_p . The coefficients for the fit and the values of E_{eff} used to calculate h_p are given in the embedded table.

that for each segment, ζ_p is approximately constant. We then analyze each segment separately starting with the final segment of the creep hold, where $A = A^0$ and $h_p = h_p^0$ are known, and work backward to the beginning of the creep hold.