



Cite this: *Soft Matter*, 2022, 18, 7612

Received 30th June 2022,
Accepted 21st September 2022

DOI: 10.1039/d2sm00884j

rsc.li/soft-matter-journal

Yield precursor in primary creep of colloidal gels†

Jae Hyung Cho *‡ and Irmgard Bischofberger *

Colloidal gels under constant moderate stress flow only after a prolonged solid-like deformation. Predicting the time-dependent yielding of the gels would facilitate control of their mechanical stability and transport, but early detectable signs of such delayed solid-to-fluid transition remain unknown. We show that the shear rate of colloidal gels under constant stress can forecast an eventual yielding during the earliest stage of deformation known as primary creep. The shear rate before failure exhibits a characteristic power-law decrease as a function of time, distinct from the linear viscoelastic response. We model this early-stage behavior as a series of uncorrelated local plastic events that are thermally activated, which illuminates the exponential dependence of the yield time on the applied stress. By revealing underlying viscoplasticity, this precursor to yield in the macroscopic shear rate provides a convenient tool to predict the yielding of a gel well in advance of its actual occurrence.

1 Introduction

Gels formed by networks of aggregated particles serve as pharmaceuticals, cosmetics, foods, and functionalized materials in novel biomedical and energy storage applications.^{1–3} The usefulness of particulate gels can be largely attributed to their mechanical versatility. They resist deformations like solids under small stresses, but flow like liquids under stresses above a certain threshold, termed the yield stress.^{4,5} A transition from solid-like deformation to fluid-like flow occurs immediately upon the application of a sufficiently large stress. When the applied stress is just above the yield stress, however, solid-like deformation can continue for a long time, until the system eventually fluidizes.^{6–18} Given how general such delayed solid-to-fluid transition is, capturing early signs of approaching yield would improve our ability to control stability^{19,20} and transport^{21,22} of these ubiquitous soft materials.

To date, most known precursors of yield are found during later phases of the solid-like deformation. Delayed yielding of particulate gels under constant stress, similar to that of numerous other systems including colloidal glasses,²³ microgel suspensions,^{24–26} polymer gels^{27,28} and harder materials,^{29,30} is preceded by three stages of creep.^{10,12,13,16–18} In the first stage, known as primary creep, that begins after a nearly instantaneous elastic response upon the application of the stress, the deformation continuously slows down as if the

system were about to stop deforming and statically support the load without yield. Under stresses larger than the yield stress, however, secondary creep ensues during which the strain rate stays approximately constant at a finite value. The strain rate then rapidly increases during tertiary creep, which results in the fracture of the gel network. Yielding, therefore, becomes predictable only after the onset of secondary creep by the rebound of the strain rate. This macroscopic signature of imminent failure concurs with microscopic bursts of irreversible structural rearrangements detected *via* dynamic light scattering.^{16,31}

In this work, we demonstrate that delayed yielding of particulate gels can be predicted by the temporal change in the rate of deformation already during *primary* creep. Using rotational rheometry, we show that gels composed of attractive colloidal particles exhibit a characteristic power-law decrease in the macroscopic shear rate with time prior to yielding. Distinct from the linear viscoelastic response, the power-law decay of the shear rate is observed when the strain falls within a range that is independent of the applied stress. The presence of this yield precursor mirrors permanent, plastic deformations during primary creep. We hence model the macroscopic behavior as a series of mesoscopic plastic events that are thermally activated and uncorrelated in space and time. Despite its simplicity, the model reproduces the rate of change in the shear rate occurring in a stress-independent strain range. Furthermore, the model enables us to infer the exponential dependence of the yielding time on the applied stress, consistent with experimental findings. We show that this exponential dependence implies that the mesoscopic plastic events in our gels involve particle-scale rearrangements. Our results indicate that the accumulation of local plastic deformations in colloidal

Department of Mechanical Engineering, Massachusetts Institute of Technology, Cambridge, MA 02139, USA. E-mail: jaehcho@mit.edu, irmgard@mit.edu

† Electronic supplementary information (ESI) available. See DOI: <https://doi.org/10.1039/d2sm00884j>

‡ Present address: Department of Mechanical Engineering, University of California, Santa Barbara, CA 93106, USA.



gels gives rise to an early yield precursor readily measurable on a macroscopic scale.

2 Experimental methods

2.1 Particle synthesis and characterization

To show the generality of the precursor to yield in primary creep, we use two types of gels that exhibit markedly different yielding mechanisms under oscillatory strains, yet a nearly identical power-law decay of the shear rate under constant stresses. One system is characterized by the storage and the loss moduli, G' and G'' respectively, that gradually decrease with the strain amplitude γ_0 after the linear regime. For the other system, by contrast, G' and G'' increase with γ_0 , which can be ascribed to stretching of force-bearing strands of particles,^{32–34} until the moduli sharply drop upon fracture, as shown in Fig. 1.

Both the strain-softening and the strain-hardening gels are composed of polystyrene-poly(*N*-isopropylacrylamide) (PS-PNIPAM) core-shell particles, synthesized by an emulsion polymerization protocol.^{35,36} For the synthesis of the particles of the strain-softening gel, we follow the protocol described in ref. 15, which is slightly modified from that of ref. 37. In a 1 L flask equipped with a stirrer, a reflex condenser, and a gas inlet, 25.02 g of *N*-isopropylacrylamide (NIPAM, Acros Organics) and 0.2008 g of the stabilizer sodium dodecyl sulfate (SDS, Sigma-Aldrich) are dissolved in 525.14 g of DI water. After the solution is bubbled with nitrogen for 30 min, 142.75 g of styrene (Sigma-Aldrich) is added, and the mixture is heated to 80 °C in nitrogen atmosphere. Then 0.3521 g of the initiator potassium persulfate (KPS, Acros Organics) dissolved in 15.00 g of DI water is added to the mixture. After 6 h, the dispersion is cooled to room temperature and cleaned through repeated centrifugation and supernatant exchange. For the synthesis of the particles of the strain-hardening gel, we follow an additional step of the seeded emulsion polymerization in ref. 37 that increases the thickness

of the PNIPAM shell, with slight modification of the ratio of the materials. For each 100 g of the particles obtained from the first part, 12.58 g of NIPAM and 0.8994 g of the crosslinker *N,N'*-methylenebis(acrylamide) (BIS, Sigma-Aldrich) are added, and the mixture is heated to 80 °C. After the addition of 0.1264 g of KPS dissolved in 9.43 g of DI water, the mixture is stirred for 4 h. The suspension is then cooled to room temperature, and cleaned by dialysis against DI water for approximately four weeks. When preparing samples at desired volume fractions, we density-match all samples using a H₂O/D₂O mixture of 52/48 v/v to prevent sedimentation. To minimize the effect of electrostatic interactions, we add 0.5 M of sodium thiocyanate (NaSCN) to screen the charges of the particles.

At temperatures T lower than the gelation temperatures T_g , the thermosensitive PNIPAM shells induce sufficiently long-ranged steric repulsions stabilizing the particles. Both types of shells, however, are negligibly thin compared to the PS cores, such that when the shells shrink with increasing T , the particles aggregate by van der Waals attraction. We estimate the gelation temperatures T_g by measuring the temperature at which the storage modulus G' becomes larger than the loss modulus G'' at a frequency $\omega = 6.3$ rad s⁻¹ during a temperature ramp experiment at a ramp rate of 0.2 °C min⁻¹, sufficiently slow to ensure uniform sample temperature.¹⁵ For the strain-softening gel and the strain-hardening gel particles, $T_g = 27.3$ °C and 25.5 °C, respectively. Since both types of particles are made of dense PS cores covered in much thinner PNIPAM shells,^{15,37} we measure the volume fraction of our samples from the changes in their mass after drying in an oven, and assuming that the particle density is equal to the density of polystyrene $\rho = 1.05$ g cm⁻³.

We perform all our experiments at $T = 30$ °C at which both types of particles aggregate by van der Waals attraction to form kinetically arrested networks of uniformly sized clusters with a fractal dimension $d_f = 1.8 \pm 0.1$,^{35,36} a signature of gels formed by diffusion-limited cluster aggregation.^{38–41} The hydrodynamic radii a measured via dynamic light scattering (BI-200SM, Brookhaven Instruments) at $T = 30$ °C, are 90.3 ± 1.6 nm and 116.3 ± 1.8 nm for the strain-softening and the strain-hardening gel particles, respectively. Since all the measurements are taken at $T = 30$ °C, the temperature dependence of the attraction strength between the particles does not play a role in our findings.

2.2 Rheometry

We use a stress-controlled rheometer (DHR-3, TA Instruments) with a cone-plate geometry of diameter 40 mm. Sandpaper (grit size 600: average diameter 16 µm) is attached to the geometry to minimize wall slip, and the edge of the loaded sample is sealed with light mineral oil (Sigma-Aldrich) to prevent evaporation. Sample loading on the rheometer is done at a lower temperature $T = 20$ °C $< T_g$ at which both types of particles are stable. Before each experiment, the sample is rejuvenated at $T = 20$ °C while being presheared at a shear rate $\dot{\gamma} = 500$ s⁻¹ for 180 s. Gelation is then initiated by rapidly increasing the temperature to $T = 30$ °C $> T_g$ at a sample time $t_s = 0$ s. This protocol allows us to prepare the sample without shear history. We let the gel

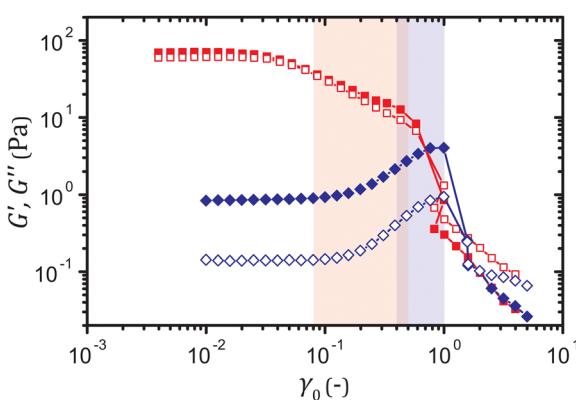


Fig. 1 Storage modulus G' (filled) and loss modulus G'' (open) as a function of the strain amplitude γ_0 at a frequency $\omega = 0.63$ rad s⁻¹ for the strain-softening (red squares) and the strain-hardening (blue diamonds) gels at a particle volume fraction $\phi = 5.0\%$. The highlighted ranges of γ_0 correspond to the strain ranges in which the yield precursor appears in creep.



evolve until $t_s = 2200$ s before applying a constant stress σ_0 , to ensure aging is slower than the deformation due to σ_0 , as described in the ESI† (Fig. S1).⁴²

3 Results

3.1 Experiments: characteristic power-law decay of the shear rate before yield

Both types of gels yield after delays when subject to stresses slightly higher than their yield thresholds. Upon the application of a stress σ_0 at time $t = 0$ s, the strain $\gamma(t)$ initially exhibits an oscillatory response induced by the coupling between the instrument inertia and the sample elasticity. The amplitude of oscillation decays more rapidly for the strain-softening gel due to its larger viscoelastic moduli,^{43,44} but for either system, the true material response soon ensues in which $\gamma(t)$ gradually increases, as shown in Fig. 2(A) and (B) for a particle volume fraction $\phi = 5.0\%$. For a stress σ_0 lower than the yield threshold, the strain $\gamma(t)$ plateaus at late times. For σ_0 higher than the threshold, $\gamma(t)$ rapidly increases marking macroscopic and catastrophic yield, once it reaches a critical value that weakly decreases with σ_0 . The yield leads to fluidization of the system after which the strain γ increases linearly with time t

(Fig. S2(A)–(D), ESI†), except for the strain-hardening gels that yield after delays longer than approximately 700 s; these systems can recover their stiffness such that γ saturates to a constant, as reported for denser colloidal gels that resolidify after yield.^{17,45} For either type of gel under a stress very close to the yield stress, yielding does not occur for the duration of the experiment even when the strain increases beyond the critical range. In such an extended deformation, aging becomes no longer negligible, as described in the ESI.† This may suggest that as particles within the network slowly migrate to form more energetically stable configurations, the toughness of the slowly deforming network is enhanced.¹⁵ Effects of gel aging during creep warrant more focused investigations and are outside the scope of this work.

The shape of the strain curves $\gamma(t)$ on log–log scales depends on the magnitude of the initial strain. The curve for the time derivative of the strain, the shear rate $\dot{\gamma}(t)$, displays the temporal change in the deformation rate unaffected by the initial strain. We find that delayed yielding, given sufficiently long primary creep, is preceded by a power-law decrease in the shear rate $\dot{\gamma}(t)$ with an exponent -0.6 , as shown in Fig. 2(C) and (D). For the strain-softening gels that do not yield, the shear rate obeys a power law $\dot{\gamma} \sim t^{-0.4}$ until it rapidly approaches zero, which reflects the viscoelastic spectrum in the inset of Fig. 2(C).

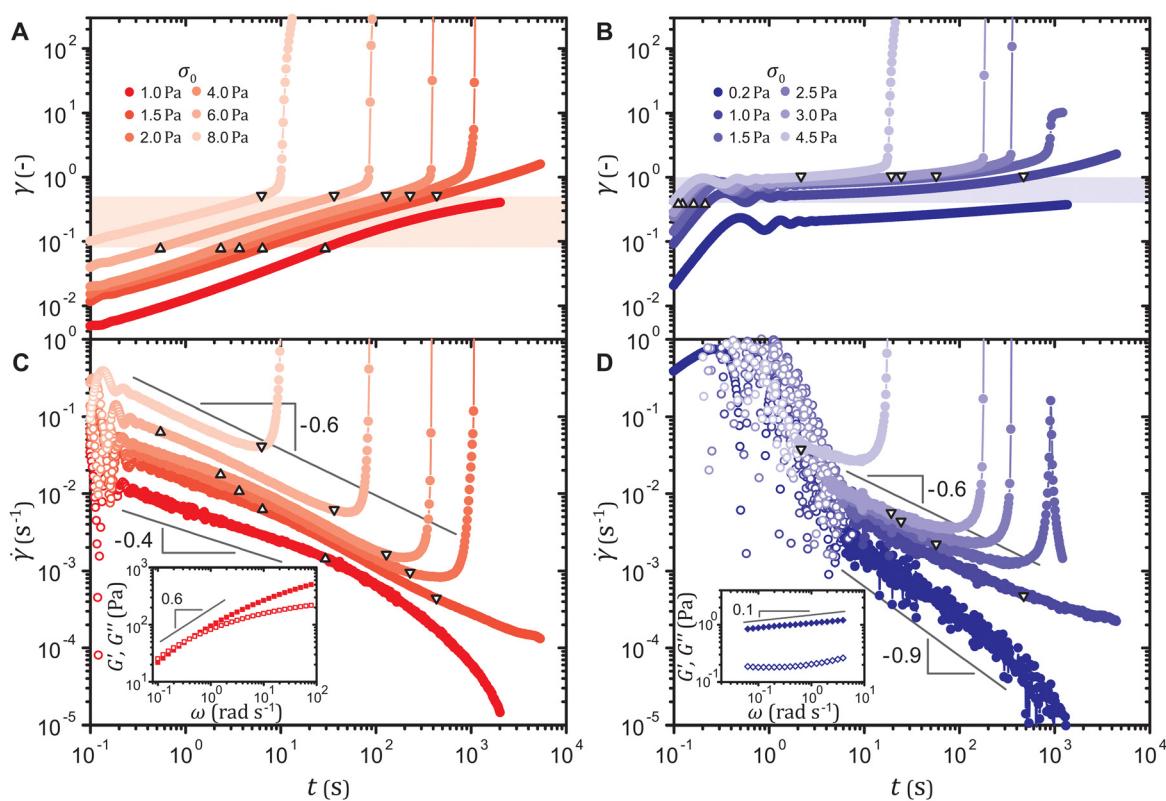


Fig. 2 Strain $\gamma(t)$ of (A) the strain-softening and (B) the strain-hardening gels at $\phi = 5.0\%$ during creep under different stresses σ_0 applied at time $t = 0$ s. Triangles and inverted triangles represent data points corresponding to the lower and the upper bounds, respectively, of the highlighted strain ranges in (A and B). Data points of $\dot{\gamma}$ at early times dominated by the instrument inertia are marked by open symbols. Insets of (C and D): linear viscoelastic spectra (G' : filled, G'' : open), obtained at strain amplitudes $\gamma_0 = 4 \times 10^{-3}$ and 8×10^{-2} for the strain-softening and the strain-hardening gel, respectively.

Linear viscoelastic behaviors in creep and in small-amplitude oscillatory shear are equivalent in that $G^*(\omega) = 1/[i\omega\hat{J}(\omega)]$, where $G^*(\omega) \equiv G'(\omega) + iG''(\omega)$ denotes the frequency ω dependent complex modulus and $\hat{J}(\omega)$ the Fourier transform of the creep compliance $J(t) \equiv \gamma(t)/\sigma_0$.⁴⁶ Substituting $J(t) = t^\alpha H(t)$, where α is a constant and $H(t)$ the unit step function, into the relation results in $G^*(\omega) = (i\omega)^\alpha/\Gamma_f(\alpha + 1) \sim \omega^\alpha$, where Γ_f denotes the Gamma function. Hence, $G^*(\omega) \sim \omega^{0.6}$ dictates that $\dot{\gamma}(t) \sim t^{0.6-1} \sim t^{-0.4}$. For the gels that do yield, the linear viscoelastic response lasts until the strain $\gamma(t)$ enters a range $\gamma = 0.08\text{--}0.5$ highlighted in Fig. 2(A), where a distinct power-law behavior $\dot{\gamma} \sim t^{-0.6}$ appears, irrespective of the applied stress σ_0 . This range of strain γ agrees with the range of strain amplitude γ_0 in which the gel strain-softens under oscillatory strains, as shown in Fig. 1, which suggests that $\dot{\gamma} \sim t^{-0.6}$ is associated with plastic deformation. The strain-hardening gels exhibit a similar behavior, as displayed in Fig. 2(D). For those that do not yield, the shear rate rapidly decreases after the linear viscoelastic response $\dot{\gamma}(t) \sim t^{0.1-1} \sim t^{-0.9}$, while for those that do yield, the characteristic power law $\dot{\gamma}(t) \sim t^{-0.6}$ appears in the strain range $\gamma = 0.4\text{--}1.0$ corresponding to the range of γ_0 in which the gel strain-hardens. For the yielding strain-hardening gels, the strain γ enters this range from the outset, and hence the transition from the linear viscoelastic response to $\dot{\gamma}(t) \sim t^{-0.6}$ is not observed. Least-squares fitting to the shear rate $\dot{\gamma}(t)$ in these strain ranges results in power-law exponents of -0.60 ± 0.05 for the strain-softening gel and -0.57 ± 0.01 for the strain-hardening gel.

This power-law decrease in the shear rate $\dot{\gamma}(t) \sim t^{-0.6}$ consistently emerges prior to delayed yielding in all our

samples, when the total strain exceeds the linear viscoelastic regime and viscoplasticity dominates primary creep. Both the strain-softening and the strain-hardening gels for volume fractions other than $\phi = 5.0\%$ exhibit $\dot{\gamma}(t) \sim t^{-0.6}$ before yielding, as shown in Fig. 3(A)–(D). Altering the linear viscoelasticity by mixing the two types of particles to form a binary gel network³⁶ also does not affect the exponent -0.6 , as displayed in Fig. 3(E) and (F). The frequency dependence of the moduli $G', G'' \sim \omega^{0.25}$ shown in the inset of Fig. 3(F) leads to $\dot{\gamma}(t) \sim t^{0.25-1} = t^{-0.75}$ in the linear viscoelastic regime, followed by the characteristic decay $\dot{\gamma}(t) \sim t^{-0.6}$ as the strain γ falls within the range $0.5\text{--}0.8$ independent of the applied stress σ_0 . We therefore identify $\dot{\gamma}(t) \sim t^{-0.6}$ as a common yield precursor in our gels during primary creep.

3.2 Model: macroscopic manifestation of accumulating local plastic events

The shear rate exhibits the characteristic power-law decay in our experiments when the total strain enters a range where plastic deformations can occur. Most yield stress fluids indeed undergo plastic deformations prior to macroscopic yielding.^{4,5} The yield precursor thus likely stems from incipient viscoplastic deformations in the gels under constant stresses. Inspired by the development of elastoplastic models,^{47–54} we build a minimal model that reproduces the temporal evolution of the shear rate $\dot{\Gamma} \sim n^{-0.6}$, where $\dot{\Gamma}$ and n denote the dimensionless shear rate and time respectively, in a stress-independent range of strain Γ through a series of local

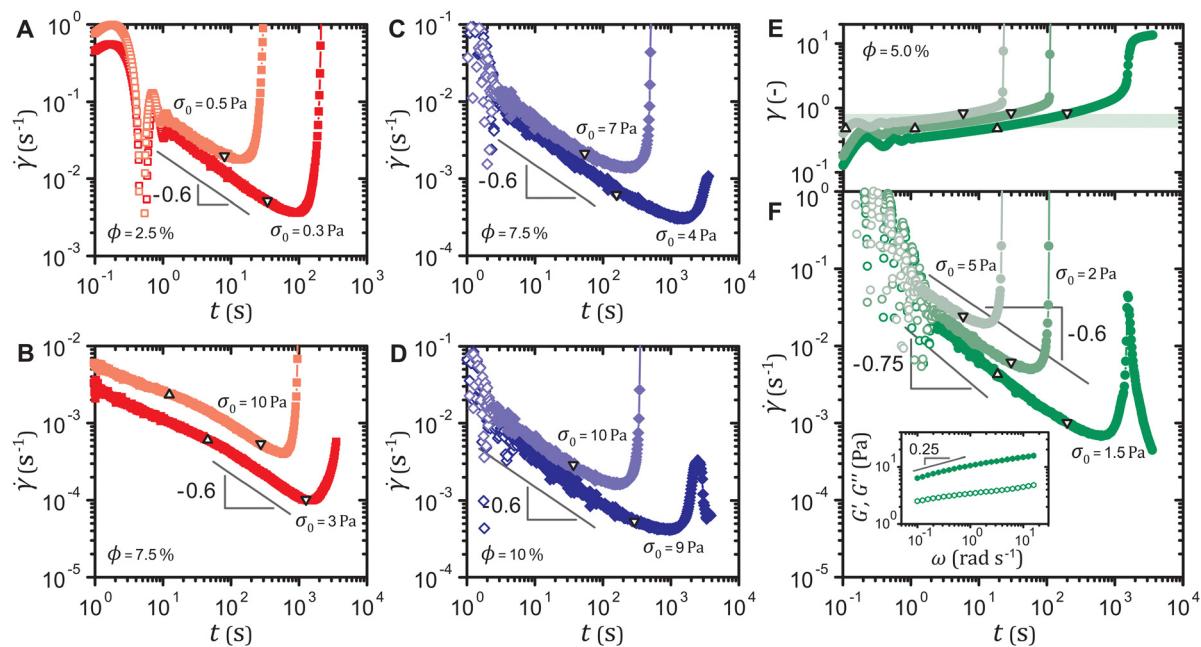


Fig. 3 Shear rate $\dot{\gamma}(t)$ of the strain-softening gel at (A) $\phi = 2.5\%$ and (B) $\phi = 7.5\%$ and the strain-hardening gel at (C) $\phi = 7.5\%$ and (D) $\phi = 10\%$. (E) Strain $\gamma(t)$ and (F) shear rate $\dot{\gamma}(t)$ of the binary gel with a volume ratio 1:1 of the strain-softening to the strain-hardening gel particles at a total particle volume fraction $\phi = 5.0\%$. Triangles and inverted triangles represent data points corresponding to the lower and the upper bounds, respectively, of a fixed strain range for each system. Inset of (F): linear viscoelastic spectrum (G' : filled, G'' : open) of the binary gel. In all cases, delayed yielding is preceded by $\dot{\gamma} \sim t^{-0.6}$ during primary creep.



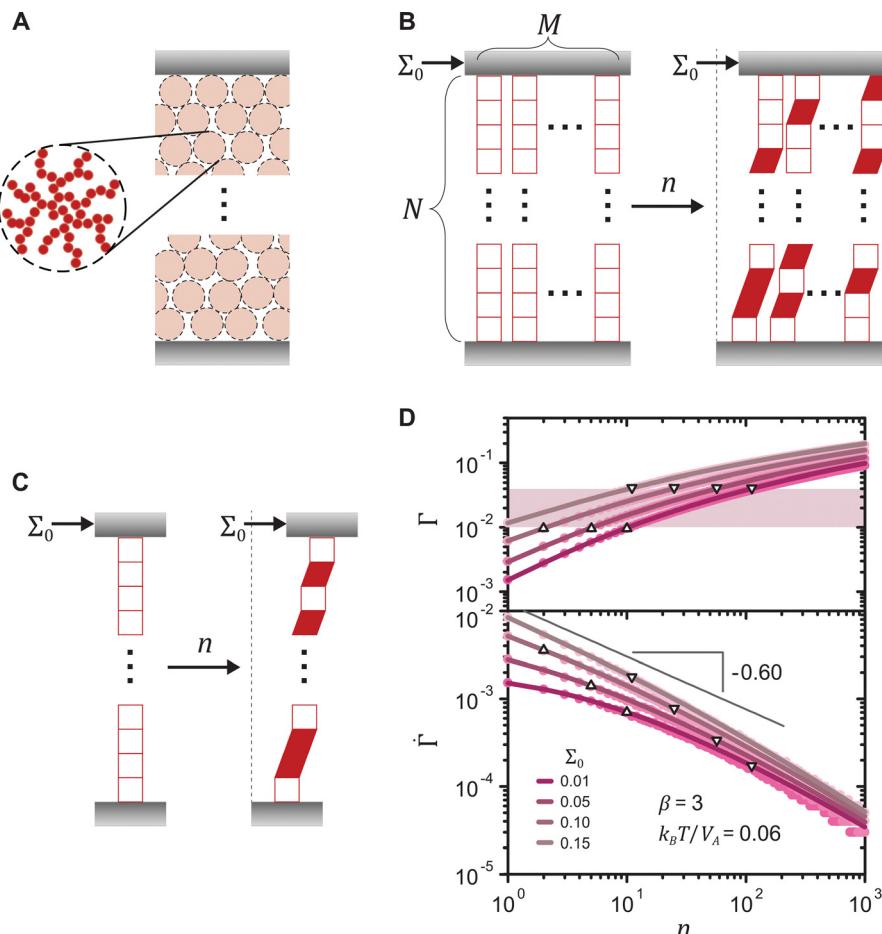


Fig. 4 (A) Schematic of the mesoscopic structure of a colloidal gel that comprises a network of uniformly sized fractal clusters between two rigid plates. (B) Coarse-graining of the gel structure to a set of M parallel arrays, each of which is formed by N serially connected mesoscopic units (open squares). Upon the application of a constant stress Σ_0 at time $n = 0$, the number of plastically deformed units (filled parallelograms) gradually increases with n . (C) One-dimensional representation of the model. (D) Strain Γ and shear rate $\dot{\Gamma}$ of the 1D model (lines) and the 2D model (circles) under different stresses for a shape parameter $\beta = 3$ of the local yield stress distribution and a thermal energy density $k_B T / V_a = 0.06$. Triangles and inverted triangles represent data points corresponding to the lower and the upper bounds, respectively, of the highlighted strain range $\Gamma = 0.01\text{--}0.04$. For the 1D model, the results are exact averaged quantities numerically evaluated by eqn (2). For the 2D model, the results are obtained from simulations for $N = 10^5$ and $M = 10^2$.

plastic events. In elastoplastic models, amorphous solids are described as networks of uniformly sized mesoscopic blocks that can locally yield.⁴⁹ The gel microstructure composed of networks of uniformly sized fractal clusters, illustrated in Fig. 4(A), entails a mesoscopic characteristic length, the size of the clusters. Given such structural hierarchy, we coarse-grain the structure of the gel network to parallel arrays of serially connected units, as shown in Fig. 4(B). Each unit, representing a cluster or its subset, bears the uniform dimensionless applied stress Σ_0 . For simplicity, we reduce the dimension of the coarse-grained system to one by considering a single array of mesoscopic units connected in series, as displayed in Fig. 4(C). Although the arrays are depicted as perfectly vertical structures between two horizontal rigid plates in Fig. 4(B) and (C), our model assumes no specific geometry of the arrays as long as the mesoscopic units are serially connected from one boundary to the other.

To define the local yield criteria, we adopt several details of the model presented in ref. 53 and 54. The local yield threshold

Σ_y of each unit is sampled from a Weibull distribution of the shape parameter β and the scale parameter $\Sigma_y^* = 1$ (Fig. S3, ESI†). The Weibull distribution, a common extreme value distribution, can arise if the local threshold is set by the strength of the weakest link within each unit.^{55,56} At time $n = 0$, each unit with a local threshold Σ_y lower than the applied stress Σ_0 yields, resulting in the local strain $\Gamma_1 = 1$. We then assume thermal activation of plastic events with the same local strain by letting each unyielded mesoscopic unit yield with probability

$$p(\Sigma_y) = \nu \exp \left[-\frac{(\Sigma_y - \Sigma_0) V_a}{k_B T} \right] \quad (1)$$

per unit time increment, where $\nu = 1$ denotes the attempt frequency, V_a the activation volume, and k_B the Boltzmann constant. To focus on the early plastic behavior, we neglect elastic deformations of individual units and elastic couplings among different units within the system, assuming no spatio-temporal correlation among plastic events. Sparse activation of



plastic events in a viscous medium is indeed expected to cause minimal correlations among local deformations in dilute colloidal gels. Each unit is allowed to yield only once at most. Microscopically, a plastic event of a mesoscopic unit may correspond to a straightening of originally tortuous force-bearing strands of particles within a cluster as hypothesized in ref. 57. For the one-dimensional (1D) model, these local yield criteria enable us to express the average macroscopic shear rate $\dot{\Gamma}$ at time $n \geq 1$ as

$$\dot{\Gamma}(n) = \Gamma_1 \int_{\Sigma_0}^{\infty} p(\Sigma_y) [1 - p(\Sigma_y)]^{n-1} f_{\text{wb}}(\Sigma_y) d\Sigma_y, \quad (2)$$

where $f_{\text{wb}} = \beta \Sigma_y^{\beta-1} \exp(-\Sigma_y^\beta)$ denotes the probability density function for the Weibull distribution of the local yield stress Σ_y (see the ESI† for the derivation). The average total strain at time $n \geq 1$ is $\Gamma(n) = \Gamma_0 + \sum_{j=1}^n \dot{\Gamma}(j)$, where $\Gamma_0 = 1 - \exp(-\Sigma_0^\beta)$ denotes the initial strain equal to the cumulative distribution function of the Weibull distribution evaluated at $\Sigma_y = \Sigma_0$.

The resulting shear rate shows a near power-law decay with a stress-independent exponent within a strain range, as experimentally found in primary creep. For a shape parameter $\beta = 3$ and a thermal energy density $k_B T/V_a = 0.06$, for instance, we observe that the shear rate follows $\dot{\Gamma} \sim n^{-0.6}$ within a strain range $\Gamma = 0.01\text{--}0.04$ for stresses from $\Sigma_0 = 0.01\text{--}0.15$, as shown in Fig. 4(D). We note that the power-law exponent of $\dot{\Gamma}(n)$ varies as β and $k_B T/V_a$ are altered. The stress independence of the exponent within a strain range, however, does not depend on the specifics of the parameters (Fig. S4(A) and (B), ESI†). The monotonic decrease in $\dot{\Gamma}(n)$ is a consequence of the gradual depletion of unyielded mesoscopic units with lower yield thresholds, and hence reflects a statistical hardening effect.^{48,54} This effect provides an explanation for why macroscopic plastic deformation slows down under constant stress and why the power-law exponent of the shear rate depends on the corresponding total strain in the experiments. Under the assumption that each unit yields once at most, the total strain at any given time represents the level of the exhaustion of unyielded units, which dictates the likely number of plastic events during the next unit time increment.

We confirm that the stress independence of the power-law exponent of the shear rate $\dot{\Gamma}(n)$ in a fixed strain range holds true for other local yield stress distributions than the Weibull distribution commonly used in elastoplastic models.⁵² Evaluation of eqn (2) using the normal distribution instead of the Weibull distribution gives rise to a power-law creep in a fixed strain range, as shown in the ESI.† The corresponding exponent depends on the choice of the mean and the standard deviation, but is largely independent of the applied stress Σ_0 (Fig. S5(A), ESI†). Using the exponential or the uniform distribution leads to a logarithmic creep $\Gamma \sim \ln(n)$, or $\dot{\Gamma} \sim n^{-1}$, for all strain Γ , as shown both numerically and analytically in the ESI† including Fig. S5(B) and (C) (ESI†). Containing only one free parameter, neither an exponential nor a uniform distribution can be used to describe a yield threshold distribution with

a peak around a finite value and a power-law exponent that changes with the strain. Despite their limitations, the one-parameter distributions successfully capture the stress independence of the exponent.

To validate the reduction of the model dimension to 1D, we simulate the creep deformation of the two-dimensional (2D) model illustrated in Fig. 4(B). In the 2D model, a uniform strain is imposed to individual arrays of serially connected mesoscopic units at every time step to reflect the uniform macroscopic strain of a sample in the cone-plate geometry in the experiment. For each time increment, after letting each unyielded unit deform with a probability determined by eqn (1), we equate the resulting mean of the total strains of individual arrays to the uniform macroscopic strain for the entire system. For the arrays whose total strains are larger than the mean, yielded units with the largest yield thresholds are then restored, while for the arrays whose total strains are smaller than the mean, unyielded units with the smallest yield thresholds are deformed to ensure that every array has the same total strain at any given time n . Though this correlation among multiple parallel arrays slightly lowers the resultant shear rate $\dot{\Gamma}$ by skewing the distribution of the local yield thresholds of unyielded units towards larger values, the overall dependence of $\dot{\Gamma}$ on time n closely agrees with the results from the 1D model, as shown in Fig. 4(D) and Fig. S4(A and B) in the ESI.†

The initial slow changes in the shear rate $\dot{\Gamma}(n)$ before the characteristic power law $\dot{\Gamma} \sim n^{-0.6}$, displayed in Fig. 4(D), cannot be observed in the experiments, in which the viscoelasticity, absent in the model, dominates the system response at low strains. Once the strain in the experiments reaches a level that requires permanent deformations, the viscoelastic response is subdued, rendering the viscoplastic response dominant as captured by the model. After the power law $\dot{\Gamma} \sim n^{-0.6}$, the accumulation of yielded units in real systems should eventually induce stronger correlations among different local plastic events leading to a proliferation of irreversible deformations throughout the gel network prior to the macroscopic yield.¹⁶ Our model, which explicitly represents the onset of plasticity by excluding such correlations, continues to display a decreasing shear rate $\dot{\Gamma}$ at late times without secondary or tertiary creep.

3.3 Exponential dependence of yield time on applied stress

In the experiments, we find that both the strain-softening and the strain-hardening gels exhibit a yield time τ_y , measured at the inflection point of the shear rate $\dot{\gamma}(t)$ for the fluidizing samples or at the local peak of $\dot{\gamma}(t)$ for the re-solidifying ones, that decreases with the stress σ_0 as $\tau_y \sim \exp(-\sigma_0/\sigma^*)$, where σ^* denotes a characteristic stress, as shown in Fig. 5(A). Such exponential dependence of τ_y on σ_0 , reported for various other colloidal gels, has been mainly attributed to the thermal activation of interparticle bond rupture.^{7,8,10–13,18}

Our model predicts the exponential stress dependence of the yield time without accounting for bond rupture, and in addition implies that the local plastic deformations during primary creep are accompanied by network rearrangements triggered at

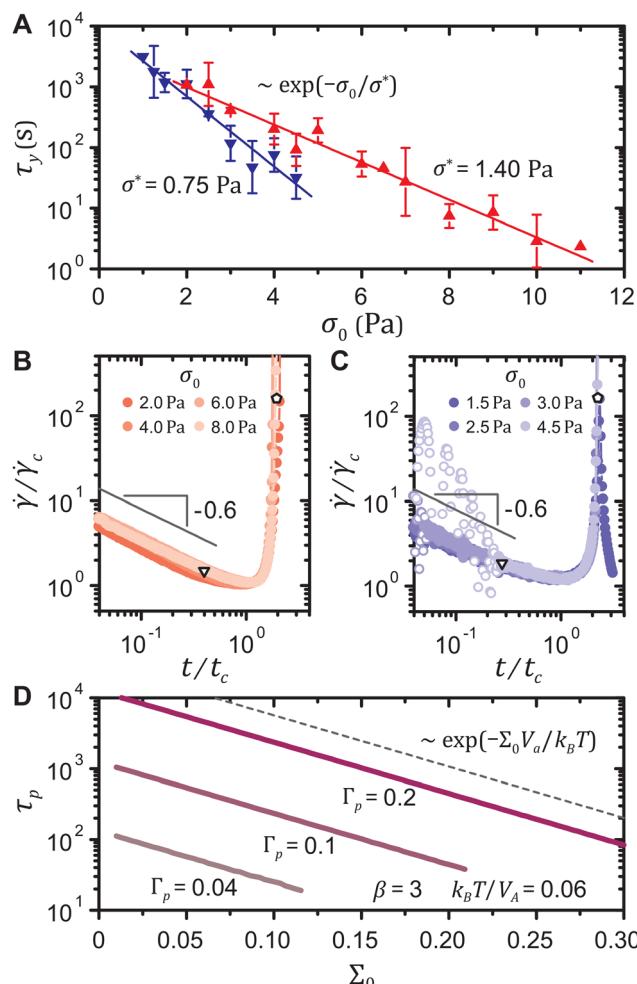


Fig. 5 (A) Experimental yield time τ_y as a function of stress σ_0 for the strain-softening (triangles) and the strain-hardening (inverted triangles) gels with exponential fits at $\phi = 5.0\%$. Data points with error bars represent averaged quantities over two to four experiments. Error bars denote the corresponding standard deviations. Shear rate $\dot{\gamma}(t)$ of (B) the strain-softening gel and (C) the strain-hardening gel scaled by the minimum shear rate $\dot{\gamma}_c$ and the corresponding time t_c for different σ_0 . Inverted triangles denote the ends of the yield precursor and pentagons denote macroscopic yield. (D) Elapsed time τ_p to reach different arbitrary total strains Γ_p in the model as a function of the stress Σ_0 . All curves scale as $\tau_p \sim \exp(-\Sigma_0 V_a / k_B T)$ (dashed line).

the particle level. The experimental shear rate curves $\dot{\gamma}(t)$ from the end of the primary creep to yield for different σ_0 can be collapsed onto a master curve on log-log scales for either type of gel, as displayed in Fig. 5(B) and (C). Thus, $\tau_y \sim \exp(-\sigma_0/\sigma^*)$ requires that the time at which the yield precursor ends also exponentially decreases with σ_0 . The model indeed shows that the time elapsed τ_p to reach an arbitrary total strain Γ_p scales as $\exp(-\Sigma_0 V_a / k_B T)$, as shown in Fig. 5(D). This scaling is robust for different thermal energy densities $k_B T / V_a$, different shape parameters β of the Weibull distribution of the local yield thresholds, and even for the three other types of distributions (normal, exponential, and uniform), as delineated in the ESI† including Fig. S6(A to C). Comparing the scaling parameters between the experiments and the model, $\sigma^* = k_B T / V_a$, we

estimate the activation radii $r_a \equiv (3V_a/4\pi)^{1/3}$ to be 89.3 ± 1.8 nm and 110.1 ± 3.8 nm for the strain-softening and the strain-hardening gel, respectively, surprisingly close to the particle hydrodynamic radii $a = 90.3 \pm 1.6$ nm and 116.3 ± 1.8 nm. This agreement hints towards particle-scale rearrangements triggering the local plastic events, reminiscent of the particle-scale plasticity observed in gels under cyclic loadings.⁵⁸

4 Discussion and conclusions

We demonstrate that during the first stage of deformation, primary creep, a characteristic power-law decay of the shear rate $\dot{\gamma}(t)$ forecasts delayed yielding of colloidal gels under constant stress. The power-law decrease in the shear rate, different from the linear viscoelastic behavior, occurs in a stress-independent strain range, which corresponds to the range of input strains that cause nonlinear stress responses in strain-controlled experiments. We interpret this yield precursor as a consequence of a series of thermally activated and spatiotemporally uncorrelated plastic deformations of mesoscopic units by building a coarse-grained model that assumes a distribution of local yield stresses and a uniformly applied external stress. Although designed to account for the early-stage viscoplastic deformation only, the model furthermore illuminates how the thermal activation of plastic events even during the first stage of creep deformation manifests itself in the exponential dependence of the yield time τ_y on the stress σ_0 . These results clarify the existence of plasticity in gels during primary creep, which has been inferred from the increase in the electrical conductivity of carbon black gels under constant stresses.⁵⁹

For all our experiments, a common yield precursor appears: a power-law exponent -0.6 of the shear rate $\dot{\gamma}(t)$, which can be reproduced by our model with a set of the Weibull shape parameter $\beta = 3$ of the local yield stress distribution and the thermal energy density $k_B T / V_a = 0.06$ within a strain range $\Gamma = 0.01\text{--}0.04$. We emphasize, however, that these results do not imply that the exponent of -0.6 is universal for colloidal gels. The primary creep deformations of other types of colloidal gels prior to yielding may be better described by different exponents, which can still be captured through our model by tuning β , $k_B T / V_a$, or even the strain range of interest. What our findings consistently indicate is that the macroscopic sign of early-stage plastic deformation arises when the level of depletion of readily deformable local units reaches a certain range. Regardless of the choice of the model parameters and even the type of yield stress distribution, the temporal decrease in the shear rate can be approximated with a power-law exponent independent of the applied stress, as long as the total strain falls within a fixed range.

The coexistence of viscoelastic and viscoplastic responses in primary creep of our colloidal gels raises the question as to why viscoelasticity alone seems to successfully account for the primary creep dynamics in other colloidal gels.^{14,16} The apparent absence of viscoplasticity may be caused by coincidental similarity between viscoelastic and viscoplastic power-law exponents.

For instance, the similarity of the two exponents -0.75 and -0.6 of the binary gel in Fig. 3(F) renders the distinction slightly less obvious than it is for the strain-softening gel in Fig. 3(B), where exponents -0.4 and -0.6 represent viscoelastic and viscoplastic responses, respectively. Also, the strain range in which viscoplasticity dominates may be narrow for systems that exhibit brittleness, in which the upper end of the linear strain regime is immediately followed by the strain at macroscopic yield. Both the strain-softening and the strain-hardening gels in this work are characterized by sufficiently broad strain ranges between the linear regime and the strain at the crossover between G' and G'' , as shown in Fig. 1, thus enabling us to observe the coexistent viscoelasticity and viscoplasticity with ease.

We note that neither the experimental nor the modeling results display the double exponential dependence of the yield time on the stress, reported for some other colloidal gels.^{10,11} The presence of the two exponential regimes in these systems is ascribed to simultaneous dissociation of multiple interparticle bonds in thick force-bearing strands, which leads to rupture of the entire strands. In fractal colloidal gels like ours, the gel network is primarily composed of thin force-bearing strands, in which rearrangements on single-particle scales alone can trigger mesoscopic and macroscopic deformations. The single-particle-scale rearrangement inferred by the agreement between the activation radii and the hydrodynamic radii of the individual particles corroborates this hypothesis.

Developing a more comprehensive description of the gels in creep prior to yielding requires the model be expanded with further details. The expression for the shear rate in eqn (2) assumes that the number of serially connected mesoscopic units N within the 1D array is sufficiently large, such that $1/N \ll \dot{\Gamma}(n)$ at any given time n , for the averaging to be physically meaningful. The experimental analog of N can be found by identifying the characteristic length scale of each mesoscopic unit and the total length of an array, which likely necessitates a direct microscopic imaging of the deforming gel. Such an investigation would provide insight into how the mesoscopic units are associated with the DLCA clusters, how the particle-scale activation triggers a mesoscale plastic event, and the topology of a deformation-governing array of mesoscopic units within the 3D system. Defining realistic correlations among distinct plastic events could elucidate how this incipient plastic behavior evolves into macroscale network rearrangements and shear localizations that may even introduce cracks during secondary and tertiary creep.⁵² The difference in the duration of these later stages of creep between the strain-softening and the strain-hardening gels, exhibited by the scaled shear rate curves in Fig. 5(B) and (C), should be accounted for in modeling the spatiotemporal correlations. It is, however, the minimal nature of our model in this work that clarifies how a slowdown of viscoplastic deformation serves as a precursor to failure. Capturing this early mechanical, macroscopic signature of yield will allow for convenient prediction of solid-to-fluid transitions, and hence enable more deft manipulation of colloidal gels in diverse settings.

Author contributions

J. H. C. and I. B. designed the research. J. H. C. performed the research and analyzed the data. J. H. C. and I. B. wrote the paper.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

We thank Thibaut Divoux and Gareth H. McKinley for helpful discussions. We acknowledge support from the MIT Research Support Committee and Kwanjeong Educational Foundation, Awards No. 16AmB02M and No. 18AmB59D.

References

- 1 J. K. Sahoo, M. A. VandenBerg and M. J. Webber, *Adv. Drug Delivery Rev.*, 2018, **127**, 185–207.
- 2 A. Z. Nelson, K. S. Schweizer, B. M. Rauzan, R. G. Nuzzo, J. Vermant and R. H. Ewoldt, *Curr. Opin. Solid State Mater. Sci.*, 2019, **23**, 100758.
- 3 S. Tagliaferri, A. Panagiotopoulos and C. Mattevi, *Mater. Adv.*, 2021, **2**, 540–563.
- 4 N. J. Balmforth, I. A. Frigaard and G. Ovarlez, *Annu. Rev. Fluid Mech.*, 2014, **46**, 121–146.
- 5 D. Bonn, M. M. Denn, L. Berthier, T. Divoux and S. Manneville, *Rev. Mod. Phys.*, 2017, **89**, 035005.
- 6 C. J. Rueb and C. F. Zukoski, *J. Rheol.*, 1997, **41**, 197–218.
- 7 V. Gopalakrishnan and C. F. Zukoski, *J. Rheol.*, 2007, **51**, 623–644.
- 8 T. Gibaud, D. Frelat and S. Manneville, *Soft Matter*, 2010, **6**, 3482–3488.
- 9 M. Laurati, S. U. Egelhaaf and G. Petekidis, *J. Rheol.*, 2011, **55**, 673–706.
- 10 J. Sprakel, S. B. Lindström, T. E. Kodger and D. A. Weitz, *Phys. Rev. Lett.*, 2011, **106**, 248303.
- 11 S. B. Lindström, T. E. Kodger, J. Sprakel and D. A. Weitz, *Soft Matter*, 2012, **8**, 3657–3664.
- 12 T. Brenner, S. Matsukawa, K. Nishinari and R. Johannsson, *J. Non-Newtonian Fluid Mech.*, 2013, **196**, 1–7.
- 13 V. Grenard, T. Divoux, N. Taberlet and S. Manneville, *Soft Matter*, 2014, **10**, 1555–1571.
- 14 M. Leocmach, C. Perge, T. Divoux and S. Manneville, *Phys. Rev. Lett.*, 2014, **113**, 038303.
- 15 D. C. E. Calzolari, I. Bischofberger, F. Nazzani and V. Trappe, *J. Rheol.*, 2017, **61**, 817–831.
- 16 S. Aime, L. Ramos and L. Cipelletti, *Proc. Natl. Acad. Sci. U. S. A.*, 2018, **115**, 3587–3592.
- 17 E. Moghimi, A. B. Schofield and G. Petekidis, *J. Phys.: Condens. Matter*, 2021, **33**, 284002.
- 18 T. E. Kusuma, P. J. Scales, R. Buscall, D. R. Lester and A. D. Stickland, *J. Rheol.*, 2021, **65**, 355–370.

- 19 T. B. J. Blijdenstein, E. van der Linden, T. van Vliet and G. A. van Aken, *Langmuir*, 2004, **20**, 11321–11328.
- 20 Y. Liang, G. Gillies, H. Patel, L. Matia-Merino, A. Ye and M. Golding, *Food Hydrocolloids*, 2014, **36**, 245–255.
- 21 C. Chang, Q. D. Nguyen and H. P. Rønning, *J. Non-Newtonian Fluid Mech.*, 1999, **87**, 127–154.
- 22 G. T. Chala, S. A. Sulaiman and A. Japper-Jaafar, *J. Non-Newtonian Fluid Mech.*, 2018, **251**, 69–87.
- 23 M. Siebenbürger, M. Ballauff and T. Voigtmann, *Phys. Rev. Lett.*, 2012, **108**, 255701.
- 24 P. H. T. Uhlherr, J. Guo, C. Tiu, X. M. Zhang, J. Z. Q. Zhou and T. N. Fang, *J. Non-Newtonian Fluid Mech.*, 2005, **125**, 101–119.
- 25 F. Caton and C. Baravian, *Rheol. Acta*, 2008, **47**, 601–607.
- 26 T. Divoux, C. Barentin and S. Manneville, *Soft Matter*, 2011, **7**, 8409–8418.
- 27 S. N. Karobi, T. L. Sun, T. Kurokawa, F. Luo, T. Nakajima, T. Nonoyama and J. P. Gong, *Macromolecules*, 2016, **49**, 5630–5636.
- 28 A. Pommella, L. Cipelletti and L. Ramos, *Phys. Rev. Lett.*, 2020, **125**, 268006.
- 29 M. C. Miguel, A. Vespignani, M. Zaiser and S. Zapperi, *Phys. Rev. Lett.*, 2002, **89**, 165501.
- 30 H. Nechad, A. Helmstetter, R. El Guerjouma and D. Sornette, *Phys. Rev. Lett.*, 2005, **94**, 045501.
- 31 L. Cipelletti, K. Martens and L. Ramos, *Soft Matter*, 2019, **16**, 82–93.
- 32 T. Gisler, R. C. Ball and D. A. Weitz, *Phys. Rev. Lett.*, 1999, **82**, 1064–1067.
- 33 J. Colombo and E. Del Gado, *J. Rheol.*, 2014, **58**, 1089–1116.
- 34 M. Bouzid and E. Del Gado, *Langmuir*, 2018, **34**, 773–781.
- 35 J. H. Cho, R. Cerbino and I. Bischofberger, *Phys. Rev. Lett.*, 2020, **124**, 088005.
- 36 J. H. Cho and I. Bischofberger, *Phys. Rev. E*, 2021, **103**, 032609.
- 37 N. Dingenaerts, C. Norhausen and M. Ballauff, *Macromolecules*, 1998, **31**, 8912–8917.
- 38 P. Meakin, *Phys. Rev. Lett.*, 1983, **51**, 1119–1122.
- 39 D. A. Weitz and M. Oliveria, *Phys. Rev. Lett.*, 1984, **52**, 1433–1436.
- 40 D. A. Weitz, J. S. Huang, M. Y. Lin and J. Sung, *Phys. Rev. Lett.*, 1984, **53**, 1657–1660.
- 41 P. van Dongen and M. H. Ernst, *Phys. Rev. Lett.*, 1985, **54**, 1396.
- 42 M. Mours and H. H. Winter, *Rheol. Acta*, 1994, **33**, 385–397.
- 43 R. H. Ewoldt and G. H. McKinley, *Rheol. Bull.*, 2007, **76**, 4–6.
- 44 C. Baravian and D. Quemada, *Rheol. Acta*, 1998, **37**, 223–233.
- 45 B. J. Landrum, W. B. Russel and R. N. Zia, *J. Rheol.*, 2016, **60**, 783–807.
- 46 J. D. Ferry, *Viscoelastic Properties of Polymers*, Wiley, New York, 3rd edn, 1980.
- 47 V. V. Bulatov and A. S. Argon, *Modell. Simul. Mater. Sci. Eng.*, 1994, **2**, 167–184.
- 48 J. C. Baret, D. Vandembroucq and S. Roux, *Phys. Rev. Lett.*, 2002, **89**, 195506.
- 49 D. Rodney, A. Tanguy and D. Vandembroucq, *Modell. Simul. Mater. Sci. Eng.*, 2011, **19**, 083001.
- 50 D. Bouttes and D. Vandembroucq, *AIP Conf. Proc.*, 2013, **1518**, 481–486.
- 51 S. Merabia and F. Detcheverry, *Europhys. Lett.*, 2016, **116**, 46003.
- 52 A. Nicolas, E. E. Ferrero, K. Martens and J. L. Barrat, *Rev. Mod. Phys.*, 2018, **90**, 45006.
- 53 D. F. Castellanos and M. Zaiser, *Phys. Rev. Lett.*, 2018, **121**, 125501.
- 54 D. F. Castellanos and M. Zaiser, *Eur. Phys. J. B*, 2019, **92**, 139.
- 55 M. J. Alava, P. K. Nukala and S. Zapperi, *Adv. Phys.*, 2006, **55**, 349–476.
- 56 Z. P. Bažant, *Proc. R. Soc. A*, 2019, **475**, 20180617.
- 57 H. K. Chan and A. Mohraz, *Phys. Rev. E*, 2012, **85**, 041403.
- 58 J. M. van Doorn, J. E. Verweij, J. Sprakler and J. van der Gucht, *Phys. Rev. Lett.*, 2018, **120**, 208005.
- 59 A. Helal, T. Divoux and G. H. McKinley, *Phys. Rev. Appl.*, 2016, **6**, 064004.