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Swelling and stress–relaxation of poly(N-isopropylacrylamide) gels in the collapsed state

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Swelling and stress–relaxation under tension for poly(N-isopropylacrylamide) (PNIPA) hydrogels have been investigated in water and in liquid paraffin at various temperatures. Initial Young's modulus is much enhanced due to the change from the swollen to collapsed state, but no difference has been observed in the initial and osmotic Poisson ratios for the gels not only in the collapsed but also in the swollen states. The enhancement of modulus originates from the introduction of physical crosslinks in the collapsed state. In liquid paraffin, the gels in the collapsed phase show large degree of stress reduction in the course of stress–relaxation due to the breakdown of the physical crosslinks. The stress–relaxation induced by swelling is observed for both phases in water, but the degree of stress reduction is larger in the collapsed phase than in the swollen phase. A comparison between experimental results and theory applicable to swelling and swelling-induced stress–relaxation is made to show that the agreement is not so good. For the stress–relaxation of the collapsed gel in water, it is suggested that the swelling couples with the breakdown of physical crosslinks at short times, and the trace of the coupling is observed even at long times. © 2002 American Institute of Physics. [DOI: 10.1063/1.1507103]

I. INTRODUCTION

Poly(N-isopropylacrylamide) (PNIPA) hydrogels have been used to investigate nature of the volume phase transition of polymer gels.^{1–5} The volume phase transition of the PNIPA gels is induced by an infinitesimal change of temperature, and a swollen state emerges at temperatures lower than the transition temperature of ca. 35 °C and a collapsed phase takes place above the transition temperature. Polymer network chains in the swollen state are solvated by the hydrophobic hydration,^{6,7} and thus are very flexible, giving the gels with rubberlike behavior. In the collapsed state the gels contain solvent inside, but the amount of solvent is very low. The network chains in the collapsed state are believed to be solvated with water molecules by hydrogen bonding. The volume phase transition of PNIPA gels originates from the change of solvation mechanism around the transition temperature. The collapsed phase may show different behavior from the swollen state due to its high polymer content, but how the collapsed phase really behaves is still unclear because of lack of information on the structure and properties of the collapsed phase. The original theory of the volume phase transition of polymer gels proposed by Tanaka *et al.*⁸ is based on the Flory-type free energy. The theory can explain the volume phase transition of charged polymer gels, but cannot be applied to PNIPA gels because they are basically neutral. A new theory has been required to describe the volume phase transition of PNIPA gels, and to build up the theory, structure and properties of the collapsed state must also be clarified.

In the previous papers,^{9,10} we have shown swelling be-

havior of poly(acrylamide) (PAAm) hydrogels under tension. The stress relaxation induced by the swelling under tension, a kind of coupling between mechanical stress and swelling (or, diffusion), was observed for the gels.⁹ When a constant strain is applied to the gels, the gels slowly re-swell to an equilibrium volume, and the applied stress (or, force) decreases with almost the same rate as the volume change to an equilibrium value. A simple theory describing the swelling-induced stress–relaxation has also been proposed.^{9,10} How mechanical stress couples with diffusion gives information on the structure, and thermodynamical and mechanical properties of polymer gels. In this study, we investigate swelling and stress–relaxation behavior of PNIPA gels in the collapsed state as well as in the swollen state. The results obtained are compared with the theory^{9,10} summarized in Sec. II. A new type of stress–relaxation occurring in the collapsed phase is modeled in terms of a rate equation.

II. THEORETICAL BACKGROUND

The change of the Gibbs free energy (ΔF) of an isotropic, cubic polymer gel, with unit volume before deformation (namely, in the reference state), under an applied tension (f) can be given by¹¹

$$\Delta F = N_s k_B T [\ln(1 - \phi) + \chi \phi] + \frac{N_c k_B T}{2} [\alpha_{\parallel}^2 + 2\alpha_{\perp}^2 - 3 - \ln \alpha_{\parallel} \alpha_{\perp}^2] - f(\alpha_{\parallel} - 1). \quad (1)$$

Here, N_s is the number of solvent molecules, and k_B , T , and ϕ are, respectively, the Boltzmann constant, the absolute

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temperature, and the polymer volume fraction, χ being the polymer-solvent interaction parameter. The quantity N_c is the number of active chains in the gel, and $\alpha_{\parallel}$ and $\alpha_{\perp}$ are the stretch ratios parallel and normal to the applied tension, respectively. From Eq. (1) swelling (or osmotic) stresses parallel and normal to the tension ($\Pi_{\parallel}$ and $\Pi_{\perp}$, respectively) can be defined by⁹

$$\begin{aligned}\Pi_{\parallel} &\equiv \frac{1}{\alpha_{\perp}^2} \frac{\partial \Delta F}{\partial \alpha_{\parallel}} \\ &= \frac{k_B T}{\nu_s} [\ln(1-\phi) + \phi + \chi \phi^2] \\ &\quad + \frac{N_c k_B T}{2 \alpha_{\perp}^2} \left(2 \alpha_{\parallel} - \frac{1}{\alpha_{\parallel}} \right) - \frac{f}{\alpha_{\perp}^2},\end{aligned}\quad (2a)$$

$$\begin{aligned}\Pi_{\perp} &\equiv \frac{1}{2 \alpha_{\parallel} \alpha_{\perp}} \frac{\partial \Delta F}{\partial \alpha_{\perp}} \\ &= \frac{k_B T}{\nu_s} [\ln(1-\phi) + \phi + \chi \phi^2] + \frac{N_c k_B T}{2 \alpha_{\parallel} \alpha_{\perp}} \left(2 \alpha_{\perp} - \frac{1}{\alpha_{\perp}} \right)\end{aligned}\quad (2b)$$

with $\nu_s = (1-\phi)\alpha_{\parallel}\alpha_{\perp}^2/N_s$, and these stresses must be zero at equilibrium. Suppose that both ϕ and f are small enough, linearized equations for Eq. (2) give to the osmotic (or, equilibrium) Poisson ratio (μ_{∞}) of 1/6 and the equilibrium stress (σ_{∞} , $\sigma_{\infty} \equiv f/\alpha_{\perp}^2$) of $7G\epsilon_{\parallel}/3$. Here, G is the shear modulus given by $G = N_c k_B T$, and $\epsilon_{\parallel}$ is the strain parallel to the tension and is related to $\alpha_{\parallel}$ as $\epsilon_{\parallel} = \alpha_{\parallel} - 1$.

To describe swelling kinetics of the gel under tension, constitutive equations as well as the equation of motion are required. These equations are applied to a small volume element in the gel specimen. Both of the total stress and strain tensors for the volume element ($\mathbf{s}$ and $\mathbf{u}$, respectively) can be divided into two components:¹⁰

$$\mathbf{s} = \mathbf{s}_{os} + \mathbf{s}_e, \quad \mathbf{u} = \mathbf{u}_{os} + \mathbf{u}_e. \quad (3)$$

Here, $\mathbf{s}_e$ and $\mathbf{u}_e$ are, respectively, the external stress and strain corresponding to the applied tension, and $\mathbf{s}_{os}$ and $\mathbf{u}_{os}$ are the osmotic stress and strain, respectively, which are induced due to the free energy change by the applied tension. When the gel is a linear elastic body, $\mathbf{s}_{os}$ is related to $\mathbf{u}_{os}$ as^{8,10}

$$\mathbf{s}_{os} = 2G[\mathbf{u}_{os} - \frac{1}{3}(\text{Tr} \mathbf{u}_{os})\mathbf{I}] + K_{os}[\text{Tr}(\mathbf{u}_{os} - \mathbf{u}_{os,\infty})]\mathbf{I}, \quad (4)$$

where $\mathbf{I}$ is the unit tensor, and K_{os} is the osmotic bulk modulus related to μ_{∞} and G as $K_{os} = 2(1 + \mu_{\infty})G/3(1 - 2\mu_{\infty})$, $\mathbf{u}_{os,\infty}$ being $\mathbf{u}_{os}$ at the long time limit. On the other hand, $\mathbf{s}_e$ and $\mathbf{u}_e$ are related with each other as¹⁰

$$\mathbf{s}_e = 2G\mathbf{u}_e - p\mathbf{I} \quad (5)$$

if Poisson's ratio as a material constant (initial Poisson's ratio, μ_0) is 1/2. The quantity p is a constant determined by a boundary condition. The quantities G and μ_0 remain constant in the course of swelling under tension and are identical to those without swelling (e.g., the measured values in air). Because polymer gels actually have μ_0 close to 1/2, the value at the incompressible limit, Eq. (5) is applicable as a

constitutive equation to polymer gels. The equation of motion for the element in the course of swelling under tension is given by^{8,10}

$$\zeta \frac{d\mathbf{v}_{os}}{dt} = \nabla \mathbf{s}_{os}. \quad (6)$$

Here, ζ is the friction coefficient between the polymer network and solvent molecules, and $\mathbf{v}_{os}$ is the displace vector corresponding to the osmotic strain $\mathbf{u}_{os}$. From Eqs. (4) and (6), the time (t) evolution of volume of the element can be described by^{9,10}

$$\frac{\partial}{\partial t} \text{Tr} \mathbf{u}_{os} = D_L \nabla^2 \text{Tr} \mathbf{u}_{os}, \quad (7a)$$

$$\frac{\partial}{\partial t} (\nabla \times \mathbf{v}_{os}) = D_T \nabla^2 (\nabla \times \mathbf{v}_{os}) \quad (7b)$$

with

$$D_L = \frac{K_{os} + (4/3)G}{\zeta}, \quad D_T = \frac{G}{\zeta}. \quad (8)$$

Equation (7a) holds for $\mathbf{u}$ as well as $\mathbf{u}_{os}$, as long as $\mu_0 = 1/2$ because $\text{Tr} \mathbf{u}_e = 0$ in this case. For the swelling of a long rectangular gel with a square cross section in the course of stress relaxation, the change of volume actually corresponds to the change in the cross-sectional area (S) and S obeys the following equation:⁹

$$\begin{aligned}S &= S_0 \left\{ 1 - \frac{128}{3\pi^4} \left[\sum_{m,n}^{\text{odd}} \frac{1}{m^2 n^2} \exp(-k_{mn}t) + \mu_{\infty} \right] \epsilon_{\parallel} \right\} \\ &(\mu_0 = 1/2, \mu_{\infty} = 1/6).\end{aligned}\quad (9)$$

Here,

$$k_{mn} = \frac{m^2 + n^2}{2\tau_L} = \frac{D_L \pi^2 (m^2 + n^2)}{a_0^2}$$

and $S_0 (= a_0^2)$ is the cross-sectional area of the gel in the reference state, and m and n are odd integers. Equation (7b) for the gel in the course of stress relaxation leads to¹⁰

$$\frac{\partial}{\partial t} u_{os,\parallel} = D_T \nabla^2 u_{os,\parallel}, \quad (10)$$

where $u_{os,\parallel}$ is the component of $\mathbf{u}_{os}$ parallel to tension. The global stress σ is given from the combination of Eq. (3b) and the solution of Eq. (10) as¹⁰

$$\sigma(t) = \frac{G\epsilon_{\parallel}}{3} \left[7 + \frac{128}{\pi^4} \sum_{m,n}^{\text{odd}} \frac{1}{m^2 n^2} \exp(-k'_{mn}t) \right] \quad (11)$$

with

$$k'_{mn} = \frac{m^2 + n^2}{2\tau_T} = \frac{D_T \pi^2 (m^2 + n^2)}{a_0^2}.$$

Equation (11) is based on the assumption that $\sigma_{\infty} = 2(1 + \mu_{\infty})G\epsilon_{\parallel} = (7/3)G\epsilon_{\parallel} (\mu_{\infty} = 1/6)$ and the initial stress (σ_0) is given by $\sigma_0 = 2(1 + \mu_0)G\epsilon_{\parallel} = 3G\epsilon_{\parallel} (\mu_0 = 1/2)$. It is also

FIG. 1. Temperature dependence of equilibrium volume (ν) of the cylindrical PNIPA gel.

noted that the ratio of τ_L to τ_T is given by $\tau_L/\tau_T = D_T/D_L = 2/5$ for $\mu_\infty = 1/6$.

III. EXPERIMENT

PNIPA gels were prepared by radical copolymerization of N-isopropylacrylamide (NIPA) and N,N'-methylenebis(acrylamide) (BIS) in distilled water. Ammonium peroxydisulfate (AP) was used as an initiator, and N, N, N', N'-tetramethylethylenediamine (TEMED) as an accelerator. The total monomer concentration (c_0) was settled to be 15 wt.%; 0.153 mol of NIPA, 2.54×10^{-3} mol of BIS, 6.13×10^{-4} mol of AP, and 240 μ l of TEMED were dissolved in 100 g of distilled water. The pregel solution was poured into a metal mold to prepare rectangular gels for mechanical tests, and the gelation was performed at 4 °C for 24 h. After gelation, the gels were removed from the mold, and were maintained in water at room temperature for 3–4 days and then in water at 65 °C for 2–3 weeks to remove residual agents. The cross section of the gel specimens was square in shape and the area was ca. 3.5 mm $\times$ 3.5 mm in the collapsed phase at the temperature equal to or higher than 40 °C and ca. 6 mm $\times$ 6 mm in the swollen phase at 30 °C.

Stress-strain and stress-relaxation behavior under uniaxial elongation was examined by using an Orientec RTM-500 tensile tester with a liquid bath. The filling solvent in the bath was water or liquid paraffin (nonsolvent for PNIPA), and the temperature of the solvent was controlled within ± 0.5 °C. Liquid paraffin was used to investigate stress-relaxation behavior without re-swelling by the applied tension. The initial length for elongation was 0.5 to 1.0 mm, and the crosshead speed was 0.5 mm/min. The magnitude of imposed strain for stress-relaxation experiments was ca. 0.1. The change of width and length between two marked points parallel to elongation were monitored by the combination of a CCD camera and a TV monitor. Initial Poisson's ration μ_0 was determined by the slope of the best fit line to the plot of $-\ln \alpha_\perp$ versus $\ln \alpha_\parallel$.

FIG. 2. Stress (σ) plotted against strain ($\epsilon_\parallel$) for the PNIPA gel in water at 40 °C.

IV. RESULTS

The phase behavior of the PNIPA gel in water was preliminarily investigated. Figure 1 shows the temperature (T) dependence of the equilibrium volume of the PNIPA gel prepared in a thin glass tube in diameter of about 200 μ m. The gel had the same monomer composition and concentration as those used for the mechanical measurements, and the volume in the figure (ν) was calculated by using the values of diameter and distance between two marked points in the axial direction. The volume of the gel decreases with increasing T , and the curve on heating agrees well with that on cooling. No discontinuity is also observed on the curves, indicating that the gel does not undergo the (discontinuous) volume phase transition. Due to the continuous change of volume, a phase boundary between the swollen and collapsed phases cannot be defined exactly for this gel, but in this paper we call the T region higher than ~ 34 °C the collapsed phase, and the region lower than that temperature, the swollen phase.

Figure 2 shows the stress (σ)–strain ($\epsilon_\parallel$) curve of the PNIPA gel in water at 40 °C. The maximum strain ($\epsilon_{\max}$) applied to the specimen was 0.5. At low strains a linear relation exists between σ and $\epsilon_\parallel$, but a downward deviation occurs for σ at large strains. Similar behavior was observed commonly for the collapsed gels at the other T in water as well as in liquid paraffin. For the gels in the swollen phase at 30 °C (both in water and liquid paraffin), only a linear region was observed on σ – $\epsilon_\parallel$ curves because the specimens failed at relatively low strains. Initial Young's modulus (E_0) for PNIPA gels estimated from the initial slope of σ – $\epsilon_\parallel$ curves for $\epsilon_{\max} = 0.1$ is listed in Table I. The values of E_0 are independent of T as well as of the type of outside solvent in the collapsed phase, and the values in the swollen phase (at 30 °C) are also independent of the type of outside solvent. Figure 3 shows a plot of $-\ln \alpha_\perp$ against $\ln \alpha_\parallel$ for the gel in water at 40 °C in the course of elongation of $\epsilon_{\max} = 0.1$. Although the data points are a little scattered, a linear relation

TABLE I. Initial Young's modulus (E_0), initial Poisson's ratio (μ_0), osmotic Poisson's ratio (μ_∞), relaxation time determined by volume change (τ_V), degree of stress reduction (r_s), and relaxation time determined by stress-relaxation (τ_σ) for PNIPA gels at various temperatures.

Solvent	Phase	$T/^\circ\text{C}$	$E_0/10^5 \text{ Pa}$	μ_0	μ_∞	$\tau_V/10^4 \text{ s}$	r_s	$\tau_\sigma/10^4 \text{ s}$
Paraffin	Swollen	30	0.063	$0.45_0 \pm 0.01$			0.13	0.9 ± 0.05
	Collapsed	40	1.5	$0.41_7 \pm 0.02$			0.30	2.6 ± 0.5
	Collapsed	60	1.6	$0.45_4 \pm 0.01$			0.29	1.9 ± 0.2
Water	Swollen	30	0.060	$0.43_3 \pm 0.02$	$0.24_6 \pm 0.01$	4.0 ± 0.3	0.25	6.1 ± 2.0
	Collapsed	40	1.5	$0.43_0 \pm 0.01$	$0.26_6 \pm 0.01$	1.3 ± 0.5	0.53	3.1 ± 0.2
	Collapsed	45	1.6	$0.45_6 \pm 0.01$	$0.20_6 \pm 0.01$	1.3 ± 0.4	0.58	3.6 ± 0.3
	Collapsed	50	1.5	$0.46_7 \pm 0.01$	$0.22_2 \pm 0.01$	1.3 ± 0.4	0.46	3.3 ± 0.2
	Collapsed	55	1.7	$0.44_5 \pm 0.01$	$0.24_4 \pm 0.01$	0.9 ± 0.3	0.55	2.6 ± 0.2
	Collapsed	60	1.7	$0.43_0 \pm 0.02$	$0.22_7 \pm 0.01$	0.8 ± 0.2	0.49	2.6 ± 0.3

is observed for the data points, and the slope of the best fit line gives $\mu_0 = 0.430$. The values of μ_0 at various temperatures are summarized again in Table I. The values remain almost constant, lying between 0.4 and 0.5, for all specimens examined, indicating that μ_0 is affected neither by the type of phase nor by the type of surrounding solvent.

In Fig. 4 the t dependence of $\epsilon_\perp$ after 10% elongation for the PNIPA gel in water at 30°C is shown. This clearly demonstrates that the gel starts to swell by the application of tension. The volume increases very steeply at short times and reaches a constant value around $t = 1 \times 10^5 \text{ s}$. The equilibrium value of $\epsilon_\perp$ ($\epsilon_{\perp,\infty}$) for this gel leads μ_∞ ($\equiv -\epsilon_{\perp,\infty}/\epsilon_{\text{max}}$) = 0.246, which is much smaller than the corresponding value of μ_0 (0.433). The similar plot for the gel in water at 40°C is shown in Fig. 5. Even in the collapsed state, the gel shows the re-swelling due to the applied tension, and μ_∞ for this gel specimen (0.266) is close to the value in the swollen state at 30°C (0.246). The values of μ_∞ at the other temperatures are tabulated in Table I, together with the above data. The values of μ_∞ are almost constant over the whole T range examined. The longest relaxation time determined from the swelling curve (τ_V), which corresponds to τ_L in Eq. (8), at various T are also summarized in

Table I. The values of τ_V in the collapsed phase appear to be weakly dependent on T , and are much smaller than that in the swollen state. The gels in liquid paraffin re-swelled neither in the swollen phase nor in the collapsed phase.

Figure 6 shows the T dependence of σ after 10% elongation in the swollen state at 30°C . In water [Fig. 6(a)], the stress decreases monotonically with increasing t and reaches an equilibrium value at about $2 \times 10^5 \text{ s}$, while the equilibration of stress occurs in a much shorter time scale in liquid paraffin [Fig. 6(b)]. The degree of stress reduction (r_s) is defined by using the initial stress (the stress just after elongation; σ_0) as

$$r_s = \frac{\sigma_0 - \sigma_\infty}{\sigma_0}. \quad (12)$$

The value of r_s is listed in Table I together with the longest relaxation time for the stress-relaxation (τ_σ) corresponding to τ_T in Eq. (8). For liquid paraffin r_s is 0.13 while that for water is 0.25. The relaxation time τ_σ is much smaller for liquid paraffin than for water. Similar plots for the gels in the collapsed phase at 40°C are shown in Fig. 7, and r_s and τ_σ

FIG. 3. A plot of $-\ln \alpha_\perp$ against $\ln \alpha_\parallel$ for the PNIPA gel in water at 40°C .

FIG. 4. Time (t) dependence of strain perpendicular to elongation ($\epsilon_\perp$) for the PNIPA gel at 30°C . The applied maximum strain for elongation is 0.108.

FIG. 5. Time (t) dependence of strain perpendicular to elongation ($\epsilon_{\perp}$) for the PNIPA gel at 40 °C. The applied maximum strain for elongation is 0.106.

are listed in Table I. Although the absolute value of r_s at 40 °C is quite larger than that at 30 °C both for water and liquid paraffin, there exists a large difference in r_s between the two outside solvents at 40 °C. However, the time profile of stress for the two specimens at 40 °C lies in the same time domain, differently from the gels in the swollen state at 30 °C. The values of r_s and τ_{σ} for the other specimens are summarized in Table I. As can be seen from the table, r_s is almost independent of T both in water and in liquid paraffin, and the values in liquid paraffin are higher than those in water. Concerning τ_{σ} , the two values for the collapsed phase in liquid paraffin are close to each other and are slightly low compared with the values in water.

V. DISCUSSION

A. Mechanical properties

As can be seen from Table I, E_0 in the collapsed phase is almost constant, resulting from the fact that the volume is almost constant in the collapsed phase above 40 °C. It is very interesting that E_0 in the collapsed phase is about 25 times higher than the values in the swollen phase. The volume decreases to ca. 1/4 of the volume in the swollen state at 30 °C (see, Fig. 1), meaning that the polymer concentration (c) becomes 4 times higher by the temperature elevation. As is well known, E_0 for swollen polymer networks scales with c as¹²

$$E_0 \sim c^{1/3}. \quad (13)$$

This scaling relation indicates that E_0 in the collapsed phase becomes higher than that at 30 °C, but the calculation based on the above scaling law shows that the degree of enhancement of E_0 is at highest 2 for the PNIPA gels, while experimental results lead the enhancement of 25 times, meaning that the increase in E_0 in the collapsed phase dominated by the other mechanism than the change of volume (or c). This

(a)

(b)

FIG. 6. Plots of stress (σ) vs time (t) for PNIPA gels at 30 °C in water (a) and in liquid paraffin (b). The applied maximum strains for elongation are 0.108 for water and 0.101 for liquid paraffin.

large enhancement of E_0 must be due to the introduction of physical crosslinks to the gel by the change of solvation from hydrophobic hydration to hydrogen bonding accompanied by the collapse, and the crosslink domain appears to be composed of a sequence or an aggregate of hydrogen bonds. An increase of entanglement density by the collapse, if any, could also enhance the apparent value of E_0 in the collapsed state. In this case σ must decrease to 1/25 of the initial value at equilibrium, but eventually the actual reduction was about 1/2 (see, r_s in Table I), suggesting that the enhancement of E_0 originates from the introduction of physical crosslinks. The fact that μ_0 depends neither on the type of phase nor on the type of outside solvent is natural because μ_0 is a material constant reflecting only the material (in this case, gel itself) properties. The values of μ_0 for the gels are rather close to

FIG. 7. Plots of stress (σ) vs time (t) for PNIPA gels at 40 °C in water (a) and in liquid paraffin (b). The applied maximum strains for elongation are 0.106 for water and 0.113 for liquid paraffin.

1/2, the value of the incompressible limit, and agree well with those reported for PAAm gels¹³ and poly(vinyl alcohol) gels.¹⁴

B. Swelling properties

The values of μ_∞ in the collapsed state was identical to that in the swollen state and remained constant independently of T , suggesting that as long as structural change (such as breakage of crosslinks) does not occur the collapsed phase obeys the same thermodynamics, for example, the Flory-type free energy expression composed of the mixing and rubber elasticity terms is also applicable to the collapsed phase, as the swollen phase, although the breakdown of physical crosslinks by the application of tension actually occurs in the collapsed phase to some extent, as will be discussed later. The network chains in the collapsed phase are flexible as in the case of swollen phase because the theory of

rubber elasticity is applicable to the polymer chains in the collapsed phase. The values of μ_∞ estimated from experiment are slightly higher than those for PAAm gels¹⁵ and also the theoretical prediction (1/6). As stated previously, the theory employed here is a linear one and is based on the assumption that polymer content is low enough, and the assumption is probably still valid for the swollen state but could not be applicable to the collapsed phase because the polymer content in the collapsed phase amounts to 60%. At least for the PNIPA gels in the collapsed state, the enhanced μ_∞ (or, reduced the swelling ratio) might originate from the high polymer content. The values of τ_V in the collapsed state showed a weak T dependence, but we think that the T dependence is only apparent because the standard deviation for τ_V is rather large (Table I), which originates from the difficulty in measurement; the values in the collapsed state must remain constant. The values of D_L in the collapsed phase calculated from τ_V and S_0 are almost comparable with that in the swollen phase, indicating that the enhancement of E_0 , or equivalently G , by the increase of c is canceled out by the increase of ζ in the collapsed phase [Eq. (8)].

C. Stress-relaxation in liquid paraffin

The stress-relaxation in liquid paraffin at 30 °C (in the swollen state) showed a small value of r_s (0.13) and the value of τ_σ was also very small. Of course, the swelling by tension does not contribute to this stress-relaxation process because the gel does not re-swell in liquid paraffin, suggesting that the mechanism of this stress-relaxation is identical to that for real elastomers, such as entanglement relaxation of dangling chains. In the collapsed phase, the stress-relaxation in liquid paraffin shows quite different behavior: τ_σ becomes larger comparable to the values in water and r_s becomes much large amounting to 0.3. In the collapsed phase, the contribution from such as entanglement relaxation to whole relaxation is negligible, and accordingly another mechanism becomes dominant. This must be due to the breakdown of crosslinks by the applied tension. As stated before, physical crosslinks are introduced in the collapsed phase. The applied tension, of course, cannot scissor the chemical crosslinks introduced in the sample preparation stage, but can break down the physical crosslinks. Actually, the downward deviation of the $\sigma - \epsilon_{||}$ curve in Fig. 1 suggests that the physical crosslinks are broken down by the applied tension. The stress-relaxation due to the breakdown of physical crosslinks, as observed for the gel in the collapsed phase in liquid paraffin, can be explained by a simple rate equation, if we assume that the annihilation rate is proportional to an extra stress, $\sigma(t) - \sigma_\infty$. Let n_c and n_p be the number densities of active (i.e., force supporting) chemical and physical crosslinks, respectively. The time dependence of n_p is determined by the following equation:

$$\frac{dn_p(t)}{dt} = -k_d[n_p(t) - n_p(\infty)], \quad (14)$$

where k_d is a rate constant for crosslink annihilation, $n_p(\infty)$ being the final value of n_p , Eq. (14) gives the relaxation modulus $G(t)$, applicable to the long time region, of the form

$$\frac{G(t)}{k_B T} \approx [n_c + n_p(\infty)] + [n_p(0) - n_p(\infty)] \exp(-k_d t). \quad (15)$$

Here, $n_p(0)$ is the initial value of n_p . The rate constant k_d is equal to $1/\tau_\sigma$ for liquid paraffin. Once a physical crosslink is broken in the course of stress-relaxation, the broken fragments will again form a new crosslink. This means the total number of physical crosslinks remains constant independently of t , but the new crosslink never participates in stress; the number of active crosslinks decreases monotonically with increasing t .

D. Stress-relaxation in water

In the swollen state at 30 °C, the gel in water showed larger values of r_s and τ_σ compared with the specimen in liquid paraffin, indicating that the stress-relaxation in water is strongly controlled by the swelling under tension. The ratio τ_V/τ_σ at 30 °C is 0.66 while the theory in Sec. II gives the value of 0.40. Although it is unclear at present why the difference occurs between the experimental and theoretical values, PAAM gels have also shown larger values of the ratio close to unity.⁹ The difference in r_s between the gels in water and liquid paraffin (0.12), which corresponds to the stress reduction originating from pure swelling, is slightly smaller than the theoretical value of r_s (0.22).

In the collapsed phase, the gel in water showed an enhanced stress reduction, larger value of r_s , compared with the gel in liquid paraffin, indicating the existence of the swelling-induced stress-relaxation even in the collapsed state. The stress-relaxation in the collapsed phase must have three origins: relaxation observed commonly for elastomers such as entanglement relaxation, breakdown of physical crosslinks, and swelling by the application of tension. The values of r_s and τ_σ in the collapsed phase are scattered (Table I), but both quantities appear to remain almost constant over the T range examined, suggesting that the percentage contribution from each origin to whole relaxation is unchanged over the T region of 40–60 °C. The long-time behavior is controlled by the latter two contributions. In this case, the long-time behavior in stress-relaxation may be described by the combination of Eq. (15) and an asymptotic form of Eq. (11); namely, by the following equation:

$$\sigma \approx \frac{k_B T \epsilon_{||}}{3} \left[7 + \frac{128}{\pi^4} \exp(-k'_{11} t) \right] \{ [n_c + n_p(\infty)] + [n_p(0) - n_p(\infty)] \exp(-k_d t) \}. \quad (16)$$

The experimental results for τ_σ in Table I indicate that $k'_{11} \cong k_d$. In the early stage of stress-relaxation of the gels in the collapsed state in water, both breakdown of physical crosslinks and swelling occur simultaneously inside the gels, and the breakdown must accelerate swelling; namely, swelling couples with the breakdown of physical crosslinks. The coupled swelling thus changes the rate of stress-relaxation

under tension. The fact that $k'_{11} \cong k_d$, which is the result in the long-time region, may be the trace of the coupling in the short-time domain. To clarify this point, more extensive and precise experiments will be needed.

VI. CONCLUSIONS

It was clearly shown from experiment that PNIPA gels in water swell by the application of tension even in the collapsed state, where the polymer content is very high. This must suggest that the collapsed phase has a network structure with flexible polymer chains just as a swollen phase. It was identified that whole stress-relaxation of PNIPA gels is composed of three contributions: relaxation observed commonly for elastomers, breakdown of physical crosslinks, and swelling-induced relaxation. An expression for the relaxation modulus originating from the breakdown of physical crosslinks, applicable to the long-time behavior of stress-relaxation of the PNIPA gels in the collapsed state in liquid paraffin, was presented. The expression was also extended to apply to the long-time stress-relaxation behavior of the collapsed gels in water, where the relaxation due to breakdown of physical crosslinks as well as swelling under tension is dominant. The results on the stress-relaxation of the PNIPA gels in the collapsed phase in water suggested that swelling couples with the breakdown of physical crosslinks in the short-time region, and the coupling affects the long-time behavior of the swelling and swelling-induced stress relaxation. A comparable theory with experimental results on the swelling and swelling-induced stress-relaxation was briefly introduced, and the comparison with experiment was made. The theory was not able to explain the experimental results on the swelling and swelling-induced stress-relaxation of the PNIPA gels in the quantitative level. A more refined model will be required to understand the entire picture of the swelling and stress-relaxation of collapsed PNIPA gels.

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