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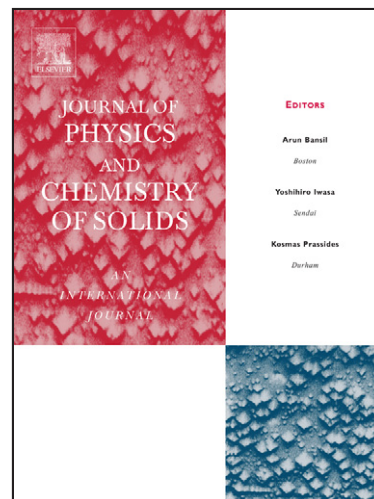
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Evaluation of the structure (or Nonlinearity) parameter x by peak-shift method from volumetric heating data of PET

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ABSTRACT

The structure parameter x , defining the temperature and structure relative contribution to the relaxation times, was evaluated applying the peak-shift method (PSM) onto volumetric heating data of poly(ethylene terephthalate) (PET) with content of crystallinity 2.6 wt. %. The method of Mercury-in-Glass Dilatometry (MIG) and three-step thermal cycle procedure were used. Two sets of heating scans were measured as a function of the amount of volume isothermal relaxation for two different relaxation temperatures below T_g (60°C and 52°C), whereas other experimental variables were kept constant. The experimentally determined heating isobar inflection temperature is a linear function of the amount of volume relaxation, which satisfy the application of PSM method for structure parameter x determination from volumetric heating data. The value of structure parameter of the used PET is independent of relaxation temperature and equal to 0.50 ± 0.02 . Finally, the shift of heating isobar inflection temperature with relaxation temperature (60°C, 56°C, 52°C and 48°C) measured for constant amount of volume isothermal relaxation was found to be linear, with the slope of 0.13.

Key words: peak-shift method, structure parameter, thermal expansion coefficient, volume relaxation, PET, glass transition temperature

INTRODUCTION

The fundamental features of amorphous solid materials, such as glass transition transformation and consecutive structural relaxation phenomenon or the behavior of relaxed structures during their isobaric heating, can be better understood and approached through the analysis of glass-formation kinetics. One of the most powerful approaches is the concept of the fictive temperature, T_f , which is used to characterize the structural state of the glass. The concept is satisfactorily expressed in an empirical relationship called Tool-Narayanaswamy-Moynihan (TNM) equation [1,2]:

$$\ln \tau_i = \ln A + \frac{x\Delta h^*}{RT} + \frac{(1-x)\Delta h^*}{RT_f} \quad (1)$$

where τ_i is the i^{th} relaxation time in the distribution of relaxation times, A means the pre-exponential factor reaching the value of τ_i at equilibrium and infinitely high temperature, Δh^* represents the apparent activation energy for structural relaxation, R is the gas constant, and x ($0 \leq x \leq 1$) stands for a non-linearity or structure parameter defining the relative contributions of temperature and structure to the relaxation times.

With the help of well designed calorimetric or volumetric experiments it is possible to determine particular parameters of TNM equation. Hutchinson *et al* developed so-called peak-shift method (PSM), which is capable, among others, to evaluate structure parameter x from calorimetric measurements [3-7]. The applicability and validity is also supported by theoretical Hutchinson's and Kovacs's work [8], where sensitiveness of parameter x to volumetric peaks is clearly demonstrated. This method is based on quantification of the peak temperature, T_p , shift at $c_p(T)$ or $\alpha(T)$ dependences caused by variation of experimental variables. The relationship between T_p and one of the variables (while the others are held constant) is usually found to be linear. The shift factor used for the determination of parameter x is calculated by linear extrapolation of the data obtained from experiments based on three-step cycles. This procedure is schematically illustrated in Fig. 1. It involves cooling of the sample at constant cooling rate, q_1 , to aging or relaxation temperature $T_a < T_g$ from annealing temperature $T_i > T_g$, where the sample was left to anneal in order to erase any previous thermal or mechanical history. It is followed by isothermal relaxation for aging time t_a . Finally, the sample is reheated at a constant heating rate, q_2 , to a temperature well above T_g .

This method can be also understood from the schematic illustration in Fig. 2. [9]. In the figure, part A clearly demonstrates an abrupt change in volume, V , which is usually measured during isobaric heating of slowly-cooled and/or annealed specimens through the glass transition region. (The same also applies to enthalpy.) This almost discontinuous increase in V with an inflection point is followed by steady state regime at sufficiently high temperature. The derivation of $V(T)$ records produces rather sharp peaks in the thermal expansion coefficient, α , which is illustrated in part B and was theoretically or experimentally shown in [8] and [9-11], or even more commonly in the specific heat capacity, c_p , [3-7] in the case of $H(T)$ dependences. The curve passes through a maximum at the mentioned peak temperature, T_p , and then decreases reaching the values characteristic of the liquid state. In Fig. 2 subscripts t_0 , t_a and ∞ define the initial volume after achieving relaxation temperature, the volume after isothermal relaxation for relaxation time t_a , and finally ∞ means equilibrium volume at T_a . Thermal expansion coefficient is mentioned as a characteristic value of glassy state, g , or liquid, l . On temperature coordinate, $T_g(q_1)$ represents conventionally determined glass transition temperature from cooling scans, T_p represents peak temperature corresponding with inflection point on $V(T)$ dependences, and T_i is initial annealing temperature before cooling.

An important role in the process is played by the used method. The signal measured on conventional calorimeters represents directly the quantity, e.g. heat flow, which can be easily re-calculated after appropriate calibration to c_p . Thus most calorimetric experimental data is more suitable for conversion of $c_p(T)$ into $H(T)$ rather than the opposite way. On the other hand, the volumetric measurements represent usually directly measured $V(T)$ data which can be converted into $\alpha(T)$.

As mentioned above, the character of the peak described by its shape, size and its temperature position is a function of experimental variables. They can be defined as q_1 , $\bar{\delta}_v$ or $\bar{\delta}_H$, q_2 and T_a . Here q_1 and q_2 mean cooling and heating rates, respectively, and quantities $\bar{\delta}_v$ or $\bar{\delta}_H$ represent the amount of volume or enthalpy isothermal relaxation at T_a . It must be kept in mind that the influence of thermal history is complex and it is necessary to separate individual effects ($q_1, \bar{\delta}_H, \bar{\delta}_v, q_2, T_a$) on T_p , so each of the effects should be treated individually.

The expansion of the peak area and peak shift to a higher temperature region is usually observed as the effect of extension of aging time, t_a [3,5-19]. Here, a linear relationship between T_p and $\bar{\delta}_H$ can be found. Hutchinson *et al* [3-9] define the normalized (dimensionless) shift by $\bar{\delta}_H$ as:

$$\delta(\bar{D}_H) = \Delta c_p \left(\frac{\partial T_p}{\partial \bar{\delta}_H} \right)_{q_1, T_a, q_2} \quad (2)$$

where $\Delta c_p = c_{p,l} + c_{p,g}$ in $[\text{J.g}^{-1}.\text{°C}^{-1}]$; $c_{p,l}$ and $c_{p,g}$ represent specific heat capacities of the liquid and glass, respectively.

It has been shown [3-7] that this shift is interrelated and depends sensitively upon x through function $F(x)$:

$$F(x) = \delta(\bar{D}_H) \quad (3)$$

Theoretically it has been found [3,4] that function $F(x)$ takes a simple hyperbolic form, Eq. (4), finally leading to the structure parameter x .

$$F(x) = x^{-1} - 1 \quad (4)$$

The reliability of this “enthalpic” approach was tested for a number of glassy polymers like PS [3], PC [6], PVAc [7], PMMA and its amorphous blends with PEO [13], poly[methyl(α -n-alkyl)acrylates] [12], for partially [14] or fully [15-17] cured epoxy resin with different curing agents, or for organic glass forming substances like glucose, fructose and their mixtures [18] or for inorganic glasses based on composition like AgI-AgPO₃-Ag₂Mo₄ [5] or recently for amorphous selenium [19].

Although this method was originally developed for both enthalpy and volumetric measurements, it has been entirely applied only onto the former, therefore dilatometric experimental data is missing in literature. The same is true for the key parameter of structural relaxation, x , for PET – it is not accessible.

Thus the main aim of this paper is to use volumetric heating data to determine the structure parameter x by the peak shift-method adapted for PET material.

EXPERIMENTAL

Tested material: Poly(ethylene terephthalate), PET, was purchased from Aldrich Chemical Company. From the original granules a specimen for dilatometric measurements was prepared by pouring the melted polymer (290°C) into room-temperature water. The density of the prepared quenched PET sample inserted into the dilatometer was measured to be $\rho = 1.3391 \text{ g.cm}^{-3}$. Comparing this value to the density of purely amorphous PET, $\rho_a = 1.335 \text{ g.cm}^{-3}$ [16] and to the calculated density of crystalline phase, $\rho_c = 1.515 \text{ g.cm}^{-3}$ [17], the degree of crystallinity can be expressed as the weight fraction, w :

$$w = \frac{\rho_c (\rho - \rho_a)}{\rho (\rho_c - \rho_a)} \quad (5)$$

The calculated value was $w = 2.6 \text{ wt. \%}$. Moreover, independent determination of sample crystallinity was carried out using a wide-angle X-ray diffraction (HZG 4 Diffractometer) giving the degree of crystallinity 2.1 % [10].

Method: A Mercury-in-Glass Dilatometer (MIG) constructed according to ASTM Standard D 864-52 [15] was used for experimental measurements, with Kovacs’s [21] remarks on the filling procedure. The concrete experimental procedure was adapted according to [24]. The dilatometer capillary was calibrated by injecting a

known amount of liquid and measuring the length of the column of liquid. Its diameter was calculated to be 0.79 ± 0.01 mm. Before sealing, the dilatometer was washed with a solution of potassium dichromate in sulfuric acid and then washed with distilled water and ethanol. The dilatometer with a specimen was filled with pure mercury (99.999%) under vacuum of about 2 Pa after its evacuation for 4 hours holding the dilatometer “bulb” with specimen at $\sim 65^\circ\text{C}$. The mercury was filtered through filter G2 before its use to remove mercury oxidizes. The amount of placed PET was approx. 1.71 cm^3 and 7.41 cm^3 of mercury. The filled dilatometer did not show any travel of mercury meniscus, at most several millimeters, after turning it upside down, i.e it was filled correctly.

All temperature programs of the dilatometer were realized by immersing the dilatometer into a precision thermostatic bath (JULABO HP-4) filled with silicon oil. The accuracy of volume changes measurements by this method was about $\sim 1 \times 10^{-5} \text{ cm}^3 \cdot \text{cm}^{-3}$ in respect to the amount of PET, mercury, dilatometer capillary bore and temperature fluctuations of thermostatic bath. The temperature lag between the temperature of silicon oil and the temperature inside the dilatometer was measured. It was determined with a K-type thermocouple placed inside the sample inserted into an additional dilatometer having the same geometry, amount of sample and mercury. For the used cooling/heating rates ($60^\circ\text{C} \cdot \text{h}^{-1}$) the lag was determined to be 0.7°C . All experimental data were corrected by this value.

Three-step thermal cycles: The above described three-step thermal cycle method, Fig. 1, was used to perform volumetric heating scans. The effect of volume isothermal relaxation, $\bar{\delta}_v$, was studied while the other variables were held constant. Also the effect of relaxation temperature for constant $\bar{\delta}_v$ was proved. The used experimental settings are listed in Table 1.

Isothermal relaxation: Isothermal volume relaxation data was collected at aging temperatures $T_a = 60^\circ\text{C}$ and 52°C . T_a was achieved by cooling from equilibrium above T_g at the cooling rate of $60^\circ\text{C} \cdot \text{h}^{-1}$. The initial time, t_0 , for collecting experimental data was the moment when T_a was reached.

RESULTS AND DISCUSSION

Volumetric glass transition temperature, T_g , is usually assessed from the volume-temperature dependence measured during sample cooling at a constant cooling rate from the equilibrium liquid state to the glassy state, and is defined as the intersection of the equilibrium liquid line with the asymptotic glassy line. Using this method, T_g of the tested PET was determined to be 68.5°C at cooling rate $q_1 = 60^\circ\text{C} \cdot \text{h}^{-1}$, as can be seen in Fig. 3 (the liquid and glassy lines are dashed). Thermal expansion coefficients were calculated as volume normalized slopes, Eq. (6), of both linear parts - above (liquid line), $\alpha_l = 5.6 \times 10^{-4} \text{ }^\circ\text{C}^{-1}$, and below (glassy line), $\alpha_g = 2.3 \times 10^{-4} \text{ }^\circ\text{C}^{-1}$, glass transition temperature, leading to $\Delta\alpha = \alpha_l - \alpha_g = 3.3 \times 10^{-4} \text{ }^\circ\text{C}^{-1}$.

$$\alpha = \frac{1}{v} \frac{dv}{dT} \quad (6)$$

Fig. 3 also includes the heating curve for the same rate $q_2 = q_1 = 60^\circ\text{C} \cdot \text{h}^{-1}$ after annealing at 56°C for 422 hours, the shape of which significantly differs from the cooling curve. A typical abrupt change can be seen on volume-temperature scan in the glass transition region. From heating scans the value of T_g can be determined as a characteristic point at the abrupt part of the heating isobar. In the present work, the inflection point, T_{ip} , was chosen for this purpose, which corresponds with the peak temperature, $T_{ip} = T_p$, on the relevant derived heating

isobaric curve. It was found that $T_{ip} = 77.3^\circ\text{C}$ from the heating mode is 8.8°C higher than T_g determined from the cooling test.

The effect of relaxation/aging time, t_a , for two different relaxation temperatures (60°C and 52°C) onto the shape and position of heating scans is presented in Fig. 4. It is nicely visible that an increase in the relaxation time leads to a decrease in specific volume with a consequent increase in the abrupt transition from the glassy to liquid line, which is followed by a shift to higher temperature. This temperature shift was consequently quantified by the slope of heating isobar inflection temperature, T_{ip} , vs. relaxation time in linear-log scale and was found to be 1.9°C .

Isothermally measured volume relaxation curves vs. relaxation time in linear-log scale for two relaxation temperatures (60°C and 52°C) and the longest experimental relaxation times (270 and 960 hours) are presented in Fig. 5. The “Volume Relaxation” is defined here as

$$v_{(t_a)} - v_{(t_0)} \quad (7)$$

where $v_{(t_0)}$ and $v_{(t_a)}$ represent specific volumes (initial and relaxation time dependent, respectively) at isothermal conditions characterized by T_a . In this figure, an initial relaxation plateau is observed at low relaxation times whereas at higher times the curves linearly decrease. Within this high relaxation time range, the intensity of volume relaxation can be quantified through volume relaxation rate (Eq. 8), which was found to be equal to $5.9 \times 10^{-4} \text{ cm}^3 \cdot \text{cm}^{-3}$ in this case.

$$\beta = \frac{1}{v} \frac{dv}{d \log t_a} \quad (8)$$

It is not difficult to show (by combination of Eqs. 3, 4, 7 and Eqs. 6, 8, 9, as given in [3]) that Hutchinson volumetric peak shift can be expressed by the following equation:

$$\hat{s}(\bar{D}_v) = \Delta \bar{\alpha} \left(\frac{\partial T_{ip}}{\partial \bar{\delta}_v} \right)_{q_1, T_a, q_2} = x^{-1} - 1 \quad (9)$$

where the “Amount of Volume Relaxation”, $\bar{\delta}_v$, is defined as $|v_{(t_a)} - v_{(t_0)}|$, the $\Delta \bar{\alpha}$ variable can be expressed as $\Delta \bar{\alpha} = \bar{\alpha}_l - \bar{\alpha}_g$, where $\bar{\alpha}_l$ and $\bar{\alpha}_g$ represent slopes (dv/dT) of the linear parts above and below T_g , respectively. In the case of the tested PET material the values were $\bar{\alpha}_l = 4.2 \times 10^{-4} \text{ cm}^3 \cdot \text{g}^{-1} \cdot ^\circ\text{C}^{-1}$ and $\bar{\alpha}_g = 1.7 \times 10^{-4} \text{ cm}^3 \cdot \text{g}^{-1} \cdot ^\circ\text{C}^{-1}$, leading to $\Delta \bar{\alpha} = 2.5 \times 10^{-4} \text{ cm}^3 \cdot \text{g}^{-1} \cdot ^\circ\text{C}^{-1}$.

In order to determine the structure parameter x from the volumetric data, it is necessary to find the slope $\Delta T_{ip} / \Delta \bar{\delta}_v$, i.e. the dependence of heating isobar inflection temperature vs. amount of volume relaxation (see Fig. 6). In the research we found slopes $4304 \text{ }^\circ\text{C} \cdot \text{g} \cdot \text{cm}^{-3}$ and $3876 \text{ }^\circ\text{C} \cdot \text{g} \cdot \text{cm}^{-3}$ for 60°C and 52°C , respectively. Then, using Eq. (9), the structure parameter x was determined for both considered temperatures as $x_{(60^\circ\text{C})} = 0.48$ and $x_{(52^\circ\text{C})} = 0.51$, i.e. equal to $x = 0.50 \pm 0.02$. In this case, the value was supposed to be relaxation temperature independent, which is in good agreement with Hutchinson *et al* [3], who proved that the non-linearity parameter x can be considered a material constant independent of all the experimental variables, such as relaxation temperature [3].

Hutchinson and co-workers in the same study [3] show that the shift of peak temperature, T_p , with relaxation temperature, T_a , for the same amount of enthalpy isothermal relaxation at different T_a can be theoretically derived to provide a value close to zero, Eq. (10):

$$\left(\frac{\partial T_p}{\partial T_a}\right)_{q_1, \bar{\delta}_V, q_2} \approx 0 \quad (10)$$

However, the volumetric experimental data from the present study suggests that the heating isobar inflection temperatures, T_{ip} , for $T_a = 60^\circ\text{C}$ are shifted to higher values by 1.5-2.1 $^\circ\text{C}$ for the same $\bar{\delta}_V$, leading to a slope of around 0.2 (from Eq. 10), as can be seen in Fig. 6. With the aim to quantify this effect much more precisely, volumetric experiments, keeping constant $\bar{\delta}_V$, were carried out for two additional relaxation temperatures (56 $^\circ\text{C}$ and 48 $^\circ\text{C}$) and summarized with previous experimental data in Fig. 7. In more detail, heating isobar inflection temperatures, T_{ip} , were determined to be 73.6 $^\circ\text{C}$ from the heating-curve after 144-hour aging at 48 $^\circ\text{C}$, 74.1 $^\circ\text{C}$ after 79 hours at 52 $^\circ\text{C}$, 74.5 $^\circ\text{C}$ after 54 hours at 56 $^\circ\text{C}$ and 75.2 $^\circ\text{C}$ after 43 hours at 60 $^\circ\text{C}$. Using this data set, the slope given by Eq. 10 was found to be 0.13 for the tested PET. This is in good agreement with enthalpy-based experimental work of Hutchinson *et al* [3], who have revealed that for polystyrene the slope varies between 0.01-0.1.

CONCLUSION

The research has proved that the peak-shift method can be applied to the volumetric data obtained from Mercury-in-Glass Dilatometry. The value of the structure parameter x for PET material was found to be relaxation temperature independent and equal to 0.50 \pm 0.02. Finally, it has been revealed that increasing relaxation temperature shifts the heating scans to a higher temperature, while other experimental conditions keep a constant amount of volume relaxation.

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Figure captions

Fig. 1. The schematic illustration of the three-step cycle method.

Fig. 2. The schematic illustration of volume, V , variation with temperature, T , (Part A) and their derivative thermal expansion coefficient, α , (Part B), during heating/cooling temperature scans [9].

Fig. 3. Dependences of specific volume on temperature during uniform cooling from equilibrium above T_g and heating after annealing at 56°C for 440 hours at the $q_1 = q_2 = 60^\circ\text{C}\cdot\text{h}^{-1}$.

Fig. 4. Dependence of specific volume on temperature as a function of the amount of isothermal relaxation, $\bar{\delta}_V$, at 60°C, part A, and 52°C, part B, prior to heating.

Fig. 5. Volume relaxation isotherms at $T_a = 60^\circ\text{C}$ and 52°C after cooling from equilibrium above T_g at cooling rate $q_1 = 60^\circ\text{C}\cdot\text{h}^{-1}$.

Fig. 6. Dependence of inflection point temperature, T_{ip} , on the amount of volume relaxation at 60°C and 52°C.

Fig. 7. Dependences of specific volume on temperature during heating after annealing at different T_a for the same amount of volume isothermal relaxation, $\bar{\delta}_V$.

Tab. 1 Ranges of experimental variables defining three-step thermal cycles.

Variable	Values used			
T_0 [°C]	85	85	85	85
t_0 [h]	0.25	0.25	0.25	0.25
q_1 [°C.h ⁻¹]	60	60	60	60
T_a [°C]	60	56	52	48
t_a [h]	4, 43, 140, 270	54, 422	17, 38, 79, 240, 960	144
q_2 [°C.h ⁻¹]	60	60	60	60

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FIG. 1

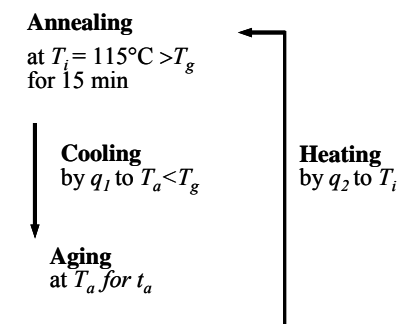


FIG. 2

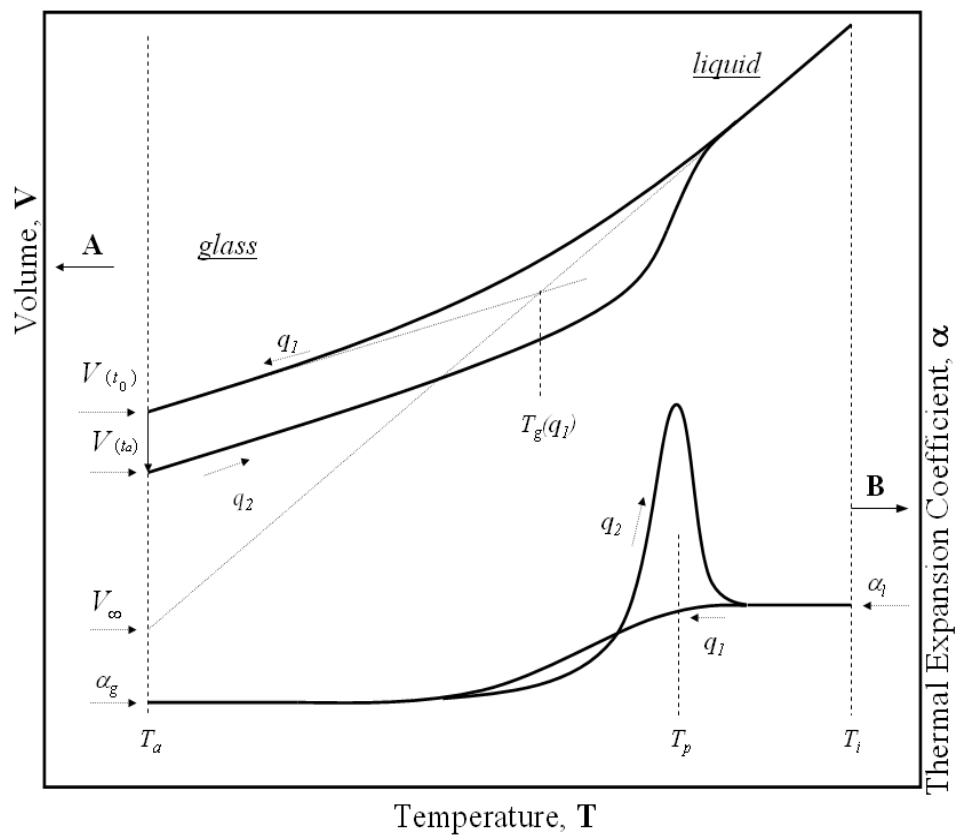


FIG. 3

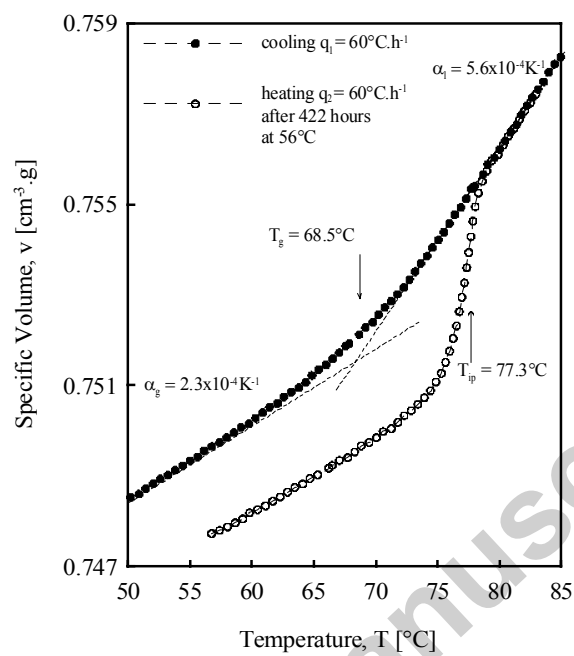


FIG. 4

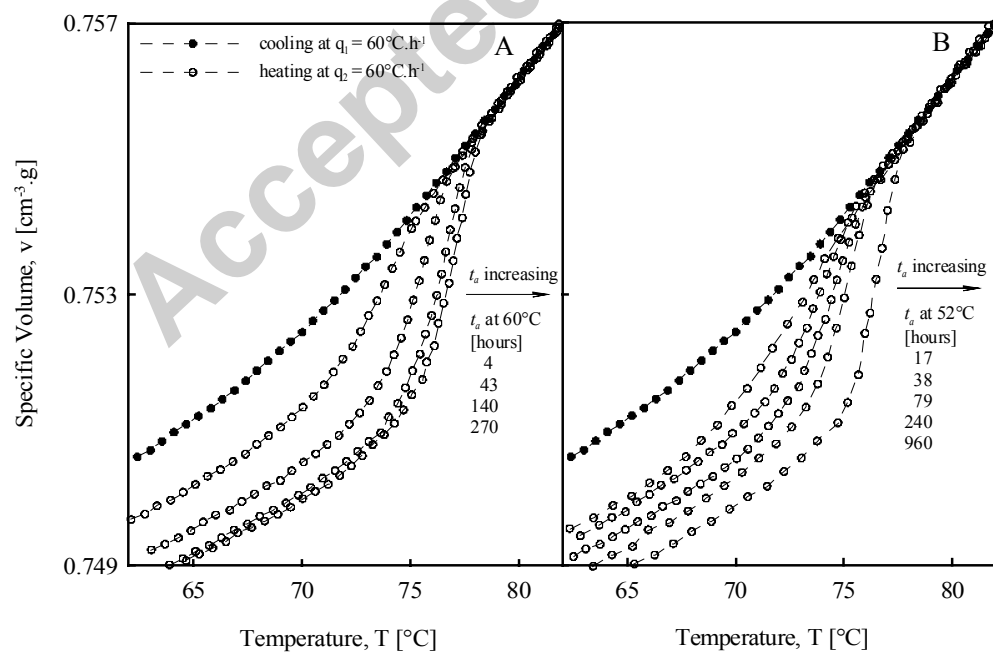


FIG. 5

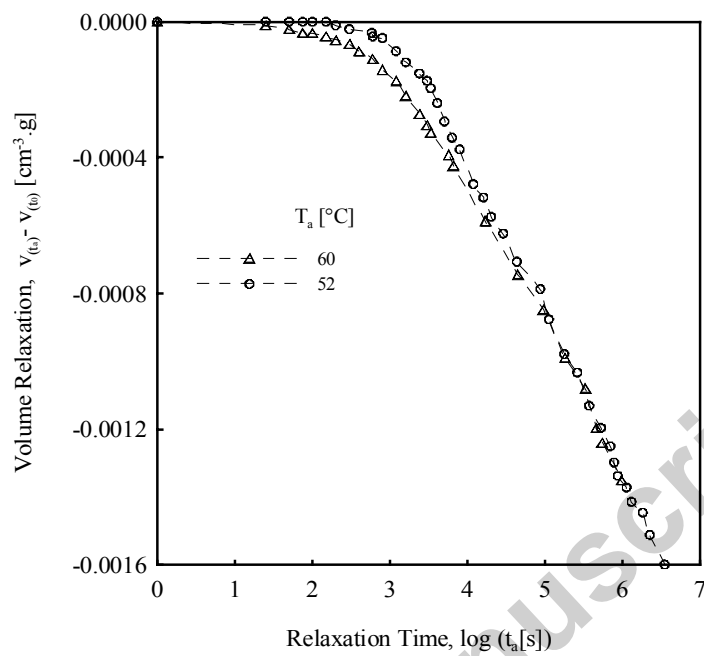


FIG. 6

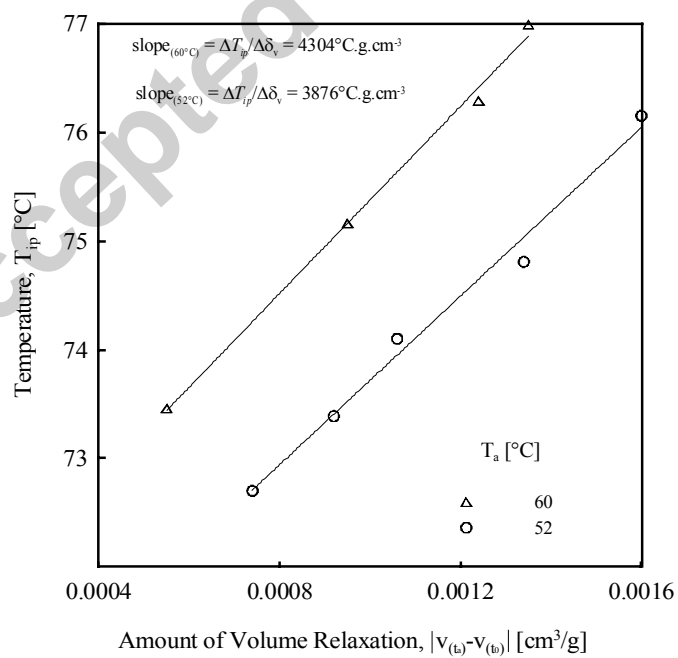


FIG. 7

