**ABSTRACT:** We examined the behavior of poly(mercaptopropyl)methylsiloxane (PMMS), characterized by a polymer chain backbone of alternate silicon and oxygen atoms substituted by a polar pendant group able to form hydrogen bonds (−SH moiety), by means of infrared (FTIR) and dielectric (BDS) spectroscopy, differential scanning calorimetry (DSC), X-ray diffraction (XRD), and rheology. We observed that the examined PMMS forms relatively efficient hydrogen bonds leading to the association of chains in the form of ordered lamellar-like hydrogen-bonded nanodomains. Moreover, the recorded mechanical and dielectric spectra revealed the presence of two relaxation processes. A direct comparison of collected data and relaxation times extracted from two experimental techniques, BDS and rheology, indicates that they monitor different types of the mobility of PMMS macromolecules. Our mechanical measurements revealed the presence of Rouse modes connected to the chain dynamics (slow process) and segmental relaxation (a faster process), whereas in the dielectric loss spectra we observed two relaxation processes related most likely to either the association–dissociation phenomenon within lamellar-like self-assemblies or the sub-Rouse mode (α′-slower process) and segmental (α-faster process) dynamics. Data presented herein allow a better understanding of the peculiar dynamical properties of polysiloxanes and associating polymers having strongly polar pendant moieties.

**INTRODUCTION**

Polysiloxanes (or silicones) are an important class of polymers having backbone consisted of alternate silicon and oxygen atoms. Because of the highly flexible structure, low chemical reactivity and toxicity, and high thermal stability, they are extensively used in, for example, adhesives, households, medicines, lubricants, textiles, and electronics industries, just to name a few. Because of this enormous versatility of applications, great effort is put into designing new enhanced siloxane-based materials. Nevertheless, it should be pointed out that this progress cannot be achieved without comprehensive studies allowing us to understand behavior and, most importantly, connect the desired properties of produced siloxane-based compounds with their structure. One can recall that the two of the most studied polysiloxanes are poly(dimethylsiloxane) (PDMS, Figure 1a) and poly(methylphenylsiloxane) (PMPS, Figure 1b). Accordingly to the classification based on the geometric dipole arrangements in polymer given by Stockmayer, both of them are considered as type B polymers, where the dipole moment is perpendicularly attached to the main chain backbone.

Recent studies of hydroxyl-terminated PDMS (assigned as HO-PDMS-OH) revealed two well-distinguishable loss peaks in the dielectric spectra of this polymer. Interestingly, a similar scenario was also noted in the mechanical data, which allowed the authors to identify the α-process (faster mode) and an additional slower one, which, by the analogy to the monohydroxy alcohols, was assigned to the mobility occurring within hydrogen-bonded supramolecular structures. Additionally, an increased ability to form self-assemblies (due to the presence of hydroxyl terminal groups in HO-PDMS-OH) leads also to a different molecular weight dependence of the glass transition temperature, $T_g$. In PDMS, $T_g$ increases with molecular weight, $M_w$, up to some limiting value of $T_g^\text{lim}$ according to the Fox–Flory relation. In the case of HO-PDMS-OH, an opposite scenario was noted. In these polymers, $T_g$ decreases with $M_w$. Nevertheless, the glass transition temperature of HO-PDMS-OH reaches a similar...
but of di-

originate from the molecular motions of comparable units,

demonstrated the presence of the two processes at first
assigned to the local segmental motion (a faster process) and
the presence of sub-Rouse modes on the low-frequency
range.17

Furth
eron correlation spectroscopy (PCS) measure-
ments performed on the low-molecular-weight PMPS also
demonstrated the presence of the two processes at first
assigned to the local segmental motion (a faster process) and
chain diffusion (slower one).20,21 However, systematic studies
indicated that although the faster process is indeed the α-
relaxation process, the slower one reflects the so-called sub-
Rouse modes of time and length scales intermediate between
the Rouse and segmental ones.22–24 Additional complement-
ary PCS experiments on PMPS and various poly(methyl-
tolylsiloxane)s (PMpTS) confirmed that both processes
originate from the molecular motions of comparable units,
but of different cooperativity.25,26 In this context, it is also
worthwhile to remind that recent studies on various PMPS,
both bulk and confined into alumina templates, also revealed
the presence of sub-Rouse modes on the low-frequency flank
of the segmental relaxation process in dielectric loss spectra.27,28
Note that no data on the presence of sub-Rouse modes within PDMS were reported so far.

Differences in the dynamics and number of relaxation modes
detected in PDMS and PMPS were discussed in terms of the
larger steric hindrance effect posed by the phenyl pendant unit
in the latter system that influenced the intermolecular cooperativity.
However, it is interesting to ask the following:
What is going to happen when the methyl side group of PDMS
is replaced by a long pendant moiety able to form the hydro-
gen bonds? Will the sub-Rouse modes be observed or
rather the Debye process? How differently will it behave in
comparison to PDMS (including HO-PDMS-OH) and PMPS?
To answer all of these questions, we studied the behavior of the
PDMS derivative, poly(mercaptopropyl)methylsiloxane (PMMS),
where the one methyl group was replaced by an alkyl pendant chain terminated with the thiol/sulfanyl (−SH)
moiety (see Figure 1c). One can mention that the ability to
form the hydrogen-bonded supramolecular structures is very
weak in the case of thiol materials. However, it is true only for
systems measured at room temperature, where these specific interactions are of comparable strength to the typical van der Waals forces.25,36 At lower temperatures, this situation is much

Figure 1. (a) Chemical structures of PDMS (a), PMPS (b), and
studied poly(mercaptopropyl)methylsiloxane, labeled as PMMS (c),
as well as its DSC thermogram (d).

value as that estimated for PDMS in the high-molecular-weight
range.17

EXPERIMENTAL SECTION

Materials. Poly(mercaptopropyl)methylsiloxane homopolymer (PMMS, 75–150 cSt) was purchased from Gelest and used as
received (molecular weight, $M_{\text{SEC}} = 2400$ g/mol, dispersity, $D = 1.26$, determined by Viscotek TDA 305 triple detection, THF as eluent).
The studied PMMS is a water-like thin liquid with the glass transition temperature $T_g = 178$ K (see Figure 1d). Note that no trace of

crystallization was observed on both cooling and heating calorimetric
scans.

Differential Scanning Calorimetry (DSC). Calorimetric measure-
ments of PMMS were performed by a Mettler-Toledo DSC apparatus equipped with a liquid nitrogen cooling accessory and an
HSS8 ceramic sensor (heat flux sensor with 120 thermocouples).
Temperature and enthalpy calibrations were performed by using indium and zinc standards. The sample was prepared in an open
aluminum crucible (40 μL) outside the DSC apparatus and measured
on heating from 150 to 298 K at a constant heating rate of 10 K/min.

Temperature-Modulated Differential Scanning Calorimetry
(TM DSC). Using a stochastic TM DSC technique implemented by a
Mettler-Toledo TOPEM , we analyzed the dynamic behavior of
PMMS in the frequency range from 4 to 30 mHz in one single
measurement at a heating rate of 0.5 K/min. In the experiment, the
temperature amplitude of the pulses of 0.5 K was selected with a
switching time range with minimum and maximum values of 15 and
30 s, respectively. The calorimetric structural relaxation times, $\tau_{\phi} = 1 / 2\pi f_{\phi}$, were determined from the temperature dependences of the real
part of the complex heat capacity, $C_p^* (T)$, obtained at different
frequencies in the glass transition region for PMMS. The glass
transition temperature $T_g$ was determined for each frequency as the

the temperature of the half-step height of $C_p^* (T)$. The

Fourier Transform Infrared Spectroscopy (FTIR). FTIR
measurements were performed with the use of a Nicolet iS50 FTIR
spectrometer (Thermo Scientific) in the frequency range from 800 to
4000 cm$^{-1}$. The bands below 800 cm$^{-1}$ were not taken into account
due to the absorption of CaF$_2$ windows. FTIR spectra were collected
with the spectral resolution of 4 cm$^{-1}$, taking 32 scans per spectrum.
The temperature experiment of FTIR measurements was regulated/
controlled through a Linkam THMS 600 heating/cooling stage
(Linkam Scientific Instruments Ltd., Surrey, UK) mounted inside the
sample stage of the IR spectrometer. The temperature changes were
measured with the cooling rate of 10 K min$^{-1}$, and the temperature
stabilization accuracy was equal to 0.1 K.

FTIR spectra of PMMS solutions in carbon tetrachloride (CCl$_4$)
were recorded by using a transmission solution cell with KBr (0.01 M)
and NaCl (0.1 M) windows with a path length of 1.03 and 0.26
mm, respectively. For each solution, 16 scans were collected in the
range from 400 to 4000 cm$^{-1}$ with a spectral resolution of 4 cm$^{-1}$.

X-ray Diffraction (XRD). The temperature-dependent wide-angle
XRD measurements were performed by using a Rigaku-Denki D/MAX RAPID II-R diffractometer with Ag K$_\alpha$ radiation (wavelength $\lambda = 0.5608$ Å), equipped with an incident beam (002) graphite monochromator and an image plate detector in the Debye–Scherrer
gometry. The sample was measured in glass capillary with a diameter
of 1.5 mm. An empty capillary was measured under the same conditions and used for background subtraction. The temperature was controlled by using an Oxford Cryostream Plus and Compact Cooler in the range from 193 to 293 K. The obtained two-dimensional diffraction patterns were converted into one-dimensional intensity data versus the scattering vector, \( q = 4\pi \sin \theta/\lambda \), where \( \theta \) is the scattering angle.

**Broadband Dielectric Spectroscopy (BDS).** Dielectric permittivity, \( \varepsilon' (\omega) = \varepsilon_r - i\varepsilon'' (\omega) \), values at ambient pressure were measured by using the impedance analyzer (Novocontrol Alpha) over a frequency range from \( 1 \times 10^{-1} \) to \( 3 \times 10^6 \) Hz. The samples were placed between two stainless-steel electrodes (diameter: 10 mm; gap: 0.05 mm) and mounted inside a cryostat. During the measurement, each sample was maintained under dry nitrogen gas flow. The temperature was controlled by a Quatro Cryosystem using a nitrogen gas cryostat, with stability better than 0.1 K. The temperature-dependent dielectric measurements were performed in the range from 173 to 303 K. It should be mentioned that for the majority of glass formers, dielectric spectra are usually shown in a permittivity, \( \varepsilon' \), representation (see Figure S5). However, to compare results obtained from mechanical and dielectric investigations, usually, latter data are converted to the modulus representation.13 Electric modulus and complex permittivity are related as follows:

\[
M^*(\omega) = \frac{1}{\varepsilon'' (\omega)} = M' (\omega) + iM'' (\omega)
\]

where real \( M' \) and imaginary \( M'' \) parts of \( M^* \) have the following forms:

\[
M' (\omega) = \frac{\varepsilon'' (\omega)}{\varepsilon'' (\omega)^2 + \varepsilon'' (\omega)^2} \quad \text{and} \quad M'' (\omega) = \frac{\varepsilon'' (\omega)}{\varepsilon'' (\omega)^2 + \varepsilon'' (\omega)^2}
\]

By use of eq 2, dielectric data were recalculated from \( \varepsilon'' (\omega) \) to \( M'' (\omega) \) as shown in Figure S3.

Collected spectra in electric modulus representation were further analyzed by the superposition of two or three Havriliak–Negami (HN) functions:

\[
M^*(\omega) = 1 + \sum_{i=1}^{2} \left[ \frac{M_{\infty} + \Delta M_i}{1 + (\omega / \omega_{HN-M_i})^{\gamma_{HN}}} \right] \exp \left[ - \left( \omega / \omega_{HN-M_i} \right)^{\gamma_{HN}} \right]
\]

where \( \omega_{HN-M} \) and \( \gamma_{HN-M} \) are the shape parameters representing the symmetric and asymmetric broadening of given relaxation peaks, \( \Delta M \) is the dielectric relaxation strength, \( \omega_{HN-M} \) is the HN relaxation time in electric modulus representation, and \( \omega \) is an angular frequency (\( \omega = 2\pi f \)).

To estimate the value of the glass transition temperature, \( T_g \), the data presented in Figure 6 were fitted by using the Vogel– Fulcher– Tamman (VFT) equation:

\[
\tau_g = \tau_{\infty} \exp \left( \frac{D_r T_g}{T_g - T_0} \right)
\]

where \( \tau_g \) is the relaxation time at finite temperature, \( D_r \) is the fragility parameter, and \( T_0 \) is the temperature, where \( \tau \) goes to infinity. To calculate the activation barrier for the \( \gamma \)-process, the Arrhenius equation was used:

\[
\tau_g = \tau_{\infty} \exp \left( \frac{\Delta E}{k_B T} \right)
\]

where \( k_B \) is the Boltzmann constant and \( \Delta E \) is the activation energy.

**Shear Rheology.** The viscosity of PMMS was determined by means of an ARES G2 rheometer. The viscosity measurement in the vicinity of the liquid–glass transition was performed by using aluminum parallel plates geometry (diameter = 8 mm). For the oscillation-frequency rheological experiments, the investigated sample was tested in the frequency range from 0.1 to 100 rad s\(^{-1}\) (12 points per decade) and over the temperature range \( T = 177–203 \) K.

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**RESULTS AND DISCUSSION**

To study the temperature evolution of the hydrogen (H)-bonding pattern in PMMS, the system was characterized by IR spectroscopy. Note that in our study we focused only on PMMS of molecular weight \( M_g = 2400 \) g/mol, which is a liquid with \( T_g = 178 \) K (Figure 1d). This is important information, considering that only in the case of low-molecular-weight polymers, similarly to HO-PDMS-OH, the formation of the supramolecular structures via H bonds was clearly observed.17 Just to add that with the elongation of the polymer backbone, the role of specific interactions becomes weaker, and the association process is being suppressed. FTIR measurements were performed from room temperature (\( T = 293 \) K) down to the liquid nitrogen temperature (\( T = 83 \) K). The evolution of the IR spectra for PMMS as a function of temperature is given in Figure 2. As observed at \( T = 293 \) K, PMMS shows the intense band located between wavenumber, \( \nu_1 \), 3000 and 2800 cm\(^{-1}\) (2957, 2927, 2881, 2862, and 2795 cm\(^{-1}\)) assigned to the asymmetric and symmetric C–H stretching vibrations of the methyl and methylene groups (\( \nu(C–H) \)). The S–H stretching vibration is found as a small peak at 2557 cm\(^{-1}\). According to the literature, the band of free (non-H-bonded) S–H group, \( \nu(S–H) \), is usually located at around 2580–2590 cm\(^{-1}\). Thus, the occurrence of the \( \nu(S–H) \) band in the PMMS spectrum at lower wavenumbers can be associated with the hydrogen bond’s formation.40,41 The Si–CH\(_3\) groups are easily recognized by a characteristic, very sharp band at 1260 cm\(^{-1}\), whereas the weaker peak appearing at 1175 cm\(^{-1}\) is assigned to the deformation vibration of the Si–CH\(_2\) groups in the propyl chain (\( \delta(Si–CH_3) \)).42 Both bands confirm that two types of alkyl groups are directly attached to a silicon atom. The origin of the bands at the region of \( \sim 900–1000 \) cm\(^{-1}\) can be associated with the C–C stretching vibrations of methylene groups.42,43 The strongest spectral feature in the IR spectrum of PMMS occurring as a broad band with an absorption maximum at 1080 cm\(^{-1}\) arises from the asymmetric Si–O–Si stretching vibrations (\( \nu_{as(Si–O–Si)} \)).42,44 As the temperature gradually decreases to \( T = 83 \) K, the intensity of IR bands increases, and they are slightly shifted to lower frequencies. The peaks associated with the C–H groups show the only small red-shift of about 2 cm\(^{-1}\) (\( \nu(C–H) \)) or 1 cm\(^{-1}\) (\( \delta(Si–CH_3) \)). Particularly interesting spectral behavior after the temperature drop is observed for the S–H stretching vibration band; that is, the \( \nu(S–H) \) band shifts to lower frequencies from 2557 to 2538 cm\(^{-1}\) (\( \Delta \nu = 19 \) cm\(^{-1}\)), and its intensity strongly increases. This red-shift is attributed to the shortening of the S–H bond length and denotes the
stronger H-bonding interactions between the S–H groups. The decreasing temperature also leads to a red-shift of the $\nu_{\text{Si-O-Si}}$ band; that is, the change of the peak frequency from 1080 to 1074 cm$^{-1}$ is observed ($\Delta \nu = 6$ cm$^{-1}$). In this case, the red-shift is associated with deformation of the PMMS structure, involving the elongation of Si–O bonds to accommodate the mercaptopropyl group linked to the inorganic silica chain. This suggests a modification of the polymer backbone conformation as the H-bonding interactions of –SH groups between neighboring chains become stronger with decreasing temperature. Complementary to the temperature-dependent studies, we also decided to perform solution-state FTIR measurement to evaluate the strength of the H bonds and tendency to the association of PMMS (see Figure S1). By use of this method, it was possible to determine the position of the SH stretching vibration peak at 2578 cm$^{-1}$ corresponding to the free (non-hydrogen-bonded) system in very diluted solutions (PMMS concentrations in CCl$_4$ < 0.01 M). As PMMS concentration increases (>0.01 M), the $\nu_{\text{SH}}$ band is red-shifted, reaching limiting value ~2557 cm$^{-1}$ for neat PMMS. This simple experiment demonstrated that PMMS forms supramolecular structures even at room temperature. Note that also the DLS measurements performed for solutions of PMMS in CCl$_4$ of various concentrations indicates a formation of small aggregates (see Figure S2).

To obtain further insight into the structure of PMMS during cooling, additional structural (XRD) studies were performed on this polymer at temperature range $T = 193–293$ K. Interestingly, the diffraction pattern of PMMS at $T = 293$ K shows two diffuse peaks in the low scattering vector range (see Figure 3a). The more intense reflection at around 1.5 Å$^{-1}$ gives average periodicity distance $d = 2\pi/Q$ of around 4.2 Å that could be associated with the nearest-neighbor distance of the short-range order between polymer macromolecules within the disordered network. On the other hand, the peak at around 0.75 Å$^{-1}$ arises due to the supramolecular cluster formation and a medium-range order with repeating distance, $d$, around 8.5 Å. Such a periodicity might correspond to the interchain separation distance when PMMS chain backbones tend to organize closely parallel to each other into lamellar-like domains (see Figure 3b).

As temperature decreases down to $T = 193$ K, a monotonic shift of both diffraction peaks toward higher Q values can be observed, suggesting a grow of polymer density and shortening of interchain distances. Interestingly, the intensity of the main diffraction peak increases, while the width sharpens with a temperature that is a typical behavior due to less thermal motions of the macromolecules, whereas the intensity of the prepeak atypically decreases with lowering the temperature. A similar scenario was observed in H-bonding monohydroxy alcohols$^{36}$ and other associating systems.$^{36}$ This effect is mostly interpreted as due to a change in the relative orientation of chains or inhomogeneous fluctuation in the density (distribution of chains in the network). In turn, the prepeak width decreases slightly with a temperature drop, and the average size of the ordered regions, $L$, increases up to 3.7 nm at $T = 193$ K. This indicates the longer-range ordering of chains due to less thermal motions of molecules and formation of the stronger H bonds via thiol moiety compared to $T = 293$ K, as found by FTIR investigations. The diffraction pattern in the wide scattering vector range showed in the inset of Figure 3a is mostly related to the intrachain structure of the studied PMMS via hydrogen bonds. As can be seen, this part of the diffraction data do not show any significant changes with decreasing temperature. Although FTIR studies indicated the elongation of Si–O bonds in the PMMS backbone with decreasing temperature, the overall structural motif of PMMS chains is preserved.

To summarize this part, one can state that presence of thiol moieties (having a weak tendency to the formation of the effective H bonds) at the end of alkyl pendant side groups induce association process that gets stronger with lowering temperature. In consequence, this phenomenon leads to the formation of lamellar supramolecular domains consisting of a few chains parallel to each other. What is more, since the examined PMMS has the H-bonding moiety (–SH group, Figure 1c) in every repeatable unit, one can expect that H bonds between neighboring chains will create some kind of cross-links leading to rather ladder-like architecture than the chain or brush-like structure discussed in the case of hydroxyl-terminated HO-PDMS-OH.$^{37}$

In the next step, we performed dielectric measurements in a wide temperature range to characterize the molecular dynamics of PMMS. Unexpectedly, it was found that there is a bimodal loss peak consisted of the two relaxation processes that tend to merge upon approaching the glass transition temperature, $T_g$ (see Figure 4). To visualize this finding in a better way, three representative dielectric loss spectra out of those presented in Figure 4a were separately shown together with the fits composed of single or superposition of the two Havriliak–Negami (HN) functions (see Figures S3 and S4). This kind of data presentation clearly indicated that the latter protocol (two HN functions) yields fits that described the obtained data much better in the whole range of frequencies when compared to the former approach. Note that the fit quality when using only one HN function was markedly worse, especially in the low frequencies (see Figure S4). This implies that the loss peak observed above $T_g$ is, in fact, a bimodal and cannot be considered as broad and asymmetric single relaxation process. Consequently, further, the collected data were analyzed with the use of at least two HN functions (see the Experimental Section). In Figure S5, we plot the HN shape parameters as well as the dielectric strength, $\Delta\varepsilon$, determined for both relaxation processes detected above $T_g$. As can be seen, $\alpha_{\text{HN}}$...
and $\beta_{\text{Rouse}}$ of the slow mode (labeled herein as $\alpha'$) are very close to the unity indicating its Debye-like nature. On the other hand, both parameters obtained for the faster (labeled as $\alpha$) relaxation are much lower than unity, suggesting that it might be the segmental process connected to the viscous flow. One can also mention that $\Delta \varepsilon$ of $\alpha'$ and $\alpha$ remains constant or slightly increases with the lowering temperature, respectively. Nevertheless, it should be mentioned that due to the proximity of both processes, there is quite large uncertainty in the determination of these parameters. Additionally, one can add that with further lowering the temperature below $T_g$, one secondary relaxation (assigned as $\gamma$) appears at high frequencies (see Figure 4b).

One can recall that usually above $T_g$ one prominent $\alpha$-relaxation process, responsible for the cooperative motions of molecules/segments, can be observed. Although in some specific molecules/polymers, this universal mobility is accompanied by the additional relaxation modes reflecting the motions of larger (sub)units. These extra modes can be considered as related to the mobility within self-assemblies (Debye process in low-molecular-weight HO-PDMS-OH⁻), motions of polymer backbone such as chain diffusion (observed in type A polymers), sub-Rouse modes (found in PMPS), or local fluctuations of some polar moieties labeled as secondary processes. Nevertheless, taking into account that generally in the case of materials with more than two OH groups per molecule Debye relaxation is barely detectable in dielectric loss spectra, it can be supposed that this kind of mobility may not be observed in PMMS that has plenty of SH moieties in one chain (Figure 1c). Moreover, it should be pointed out that the chain dynamics is not dielectrically active for polysiloxanes (considered as type B polymers), which excludes the chain diffusion origin. One can also mention that the sub-Rouse modes are rarely observed by the BDS technique. Hence, taking into account the discussion mentioned above and the puzzling nature of the additional dielectric relaxation process observed above $T_g$ we have performed comprehensive complementary mechanical measurements.

Master curves of measured storage, $G'$, and loss, $G''$, modulus of examined PMMS are shown in Figure 5. The presented master curve was constructed from the frequency dependencies of $G'$ and $G''$ measured within the range of 0.1–100 rad s⁻¹ at various temperatures (Figure S7), which were shifted horizontally by the shift factor, $\Delta T$, to superimpose at the reference temperature, $T_{\text{ref}} = 180$ K. As observed, two prominent processes can be detected, the faster (at higher frequencies) reflecting the segmental motions and an additional one at lower frequencies (Figure 5). Both modes are separated by ~4 orders of magnitude on the time scale. One can add that a similar situation, with the same difference in relaxation rates, was also reported for various PDMS characterized by different chain ends. In these polymers, the faster and the slower processes were assigned to the segmental motions and Rouse modes (related to the terminal/chain relaxation), respectively. Therefore, by the analogy, this interpretation of $G'$ and $G''$ spectra of PMMS will be adopted herein. Moreover, a direct comparison of the dielectric and rheological data presented in Figures 4 and 5 revealed that the chain mode does not correspond to any of relaxation processes detected in loss spectra (Figure 4). Moreover, when we compare the loss modulus, $G''$, spectra with the dielectric modulus, $M''$, near $T_g$ one can see that the $\alpha$-peak recorded in mechanical measurements has much different shape than the one detected in dielectric spectra (see Figure S8). Both findings discussed above indicate that actually both applied techniques monitor a different kind of mobility. Note that to set together the results obtained from the mechanical and dielectric measurements, generally loss $G''$ and dielectric $M''$ are compared (see the Experimental Section).

To explore in more detail the molecular dynamics and the origin of the two dielectric relaxation processes detected in the examined PMMS above $T_g$ we determined and analyzed temperature evolution of their relaxation times, $\tau$ (see Figure 6). Note that dielectric $\tau$ was estimated from the fitting of collected spectra in the electric modulus representation to the two (or three for the spectra measured close to $T_g$ where the contribution from the $\gamma$-process was significant) HN functions (see Figures S5 and S6). On the other hand, $\tau_g$ from the mechanical measurements were calculated by the following equation: $\tau_g = \omega^{-\nu}$, where $\omega$ is a frequency of the $G''(\omega)$ maximum. In the temperature range where the $G''$ peaks are not visible, the time–temperature superposition (TTS) rule can be applied to determine $\tau_g$. According to this criterion, $G'$
Figure 6. Temperature dependences of relaxation times obtained from dielectric, mechanical, and calorimetric measurements. Red solid and dashed lines represent the best fits to the VFT and Arrhenius equations, respectively (see the Experimental Section).

and $G''$ spectra collected at various temperature conditions form the master curve when shifted to the reference temperature $T_{ref} = 180$ K (see Figure 6) and $\tau_s$ is taken as the shift factor ($\alpha_T$). The relaxation time of the slower Rouse modes, $\tau_R$, was estimated by the approach proposed by Patkowski et al., from the crossover of the two straight lines, representing the dependencies of $\log(G') \sim 2 \log(\omega)$ and $\log(G'') \sim \log(\omega)$ in the low frequency. Then, using the relation $\log(\tau(T)) = \log(\tau(T_{ref})) + \log(\alpha_T)$, we obtained their temperature dependences. The obtained value of $\tau_s$ and $\tau_R$ were added to Figure 6. Additionally, we also performed temperature-modulated (TM) DSC measurements to determine values of $\tau_s$ near $T_g$ (see the Experimental Section), which are also included in Figure 6.

Interestingly, it was found that the relaxation times of the dielectrically active slower process agree well with the $\tau_s(T)$ dependence determined from both TM-DSC and rheological measurements, which might suggest that this process is the segmental ($\alpha$) relaxation related to the glass transition phenomenon. However, this implies that the faster mode is a secondary relaxation. This interpretation is consistent with the data reported for different kinds of polymers, including polymethacrylates. However, as one can recall generally, the $\alpha$ and secondary $\beta$ processes merge at high temperatures and tend to separate upon approaching $T_g$. Moreover, it should be highlighted that the secondary processes are characterized by the Arrhenius-like temperature dependences of relaxation times. These features indicated that the faster process could not be a secondary process. Considering this discussion and the shape parameters determined for both modes, we propose to assign the faster process as the real segmental ($\alpha$) relaxation related to the glass transition phenomenon and the slower dielectric process (labeled as $\alpha'$) to be connected to sub-Rouse mobility of larger than segmental (sub)units or the association–dissociation process of the formed supramolecular lamellar-like structures as revealed by XRD and FTIR investigations. Note that the terminal relaxation observed in the mechanical measurements and dielectric $\alpha'$ process are found to differ by 2–3 orders of magnitude in time, which exclude the chain diffusion origin of this process (see Figure 6).

For a deeper understanding of this problem, we compared temperature dependencies of the relaxation times obtained for PMMS with two different PDMS able to form H bonds (hydroxyl-terminated HO-PDMS-OH) and the van der Waals polymer (silanol-terminated PDMS). In the case of oligomeric HO-PDMS-OH ($M_n < 1100$ g/mol), the dielectric loss spectra are dominated by the slower (Debye) relaxation connected to the motions within supramolecular self-assemblies or association phenomena. However, the contribution of this process to the loss spectra decreases with $M_n$ and completely vanishes for polymers with $M_n > 2000$ g/mol. Note that data for HO-PDMS-OH were taken from ref 17. It should also be noted that both processes observed for HO-PDMS-OH merge when approaching $T_g$ similarly as in the case of studied PMMS (see Figure S9a). Moreover, the time scale separations between $\alpha'$ (or Debye) and $\alpha$ in both polymers are similar. These similarities might indicate that the slow mode in the dielectric loss spectra of PMMS could be related to the mobility within hydrogen-bonded supramolecular structures. In this context, it is worthwhile to remind that HN parameters describing the shape of the $\alpha'$ mode are very close to one that is characteristic for a Debye process ($\alpha \sim 0.9$ and $\beta \sim 0.8$). Nevertheless, one can emphasize that the Debye-like process observed in PMMS has a significantly lower amplitude than that detected in HO-PDMS-OH. This finding might be connected to different geometry and a large number of $-$SH moieties capable of forming H bonds that hold local molecular ordering via these specific interactions. PMMS is also characterized by much higher molecular weight ($M_n = 2400$ g/mol) than the mentioned HO-PDMS-OH ($M_n < 1100$ g/mol), which also influences differences between both materials. Assuming that indeed the $\alpha'$ relaxation is related to the association process leading to the formation of hydrogen-bonded (ladder-like) structure, we calculated the activation energy of associations, $E_{assoc}$, from dielectric data. For details, see refs 56 and 57. The estimated value of $E_{assoc}$ is equal to $E_{assoc} = 18$ kJ/mol. Interestingly, this energy is comparable to the net interaction energy $E \sim 10–16$ kJ/mol determined for mercaptobenzoic acids having a thiol moiety.18 The discussion mentioned above suggests that the recorded additional dielectric relaxation is a Debye process reflecting the formation of H bonds and supramolecular structure in PMMS as indicated by complementary FTIR and XRD investigations. Briefly, one can stress that generally for strongly associating systems, apart from the $\alpha$-process responsible for the vitrification phenomenon, an additional slower mode of still unknown origin (generally of exponential shape) is observed.59 However, it is not the strict rule since there are many exceptions such as metaloludine, water, the phenyl derivative of ibuprofen, and so on.60,61 In our case, this local ordering might indicate the formation of a hydrogen-bonded (ladder-like) structure of the length scale higher than segments. Considering the geometry of the nanoaggregates presented in Figure 3b, one can postulate that net dipole moment in such situation cancels out. However, it should be mentioned that the ongoing association–dissociation at $T_g$ leads to the change in dipole moments resulting in the rise to the $\alpha'$-process.60,62

Additionally, we examined the dynamics of native PDMS (of $M_n = 3700$ g/mol). Note that the measurements were performed after careful removal of monomer wastes from the purchased material, which contains a significant amount of impurities. Details concerning the purification procedure and the gel permeation chromatography (GPC) data recorded
before and after the performed actions are presented in Figure S9. Surprisingly, the dielectric loss spectra of purified PDMS measured above $T_g$ also revealed the presence of the two relaxation processes (see Figure S10b). One can recall that the recent papers devoted to the dynamics of various PDMS derivatives, including PMPs\textsuperscript{27,28} and PMPTS\textsuperscript{35} pointed out the presence of the dielectric relaxation process characterized by intermediate time and length scale between chain (Rouse modes) and segmental relaxations, assigned as sub-Rouse modes. Hence, by the analogy to these papers, the slower relaxation can be assigned as the sub-Rouse process, while the faster process is the segmental one in native PDMS. This agrees when we compare the $\tau_g$ determined from calorimetry ($T_g = 148$ K) as shown in Figure S11, the time-scale separation between sub-Rouse modes and segmental relaxation in the case of measured PDMS of $M_n = 3700$ g/mol is quite significant. Moreover, both processes do not merge upon decreasing temperature and have a significantly different amplitude with respect to the ones detected in PMMS (see Figure S10a). Nevertheless, even considering results of mechanical measurements and similarities and differences between dynamical behavior of the relaxation processes in PMMS and hydroxyl-terminated and native PDMS, we are not able to unquestionably state whether $\alpha'$ is connected to the motions of either sub-Rouse modes or the association process leading to the formation of medium-range ordered supramolecular lamella-like hydrogen-bonded nanodomains. On the other hand, the faster mode can be assigned as segmental relaxation in PMMS—especially when we take into account that $T_g$ ($T_{g,BDS} = 177$ K) determined from the dielectric data using the VFT equation (eq 4) agrees well with the one determined from the calorimetric measurements ($T_{g,DSC} = 178$ K, Figure 1b) and shape parameters of this mode are much different than unity.

As a final point of our investigations, we have also calculated the activation energy, $E_a$, of the $\gamma$-process in the glassy state. For that purpose, we fitted $\tau_g$ presented in Figure 6 to the Arrhenius equation (eq 5). The calculated activation energy reaches $E_a = 25$ kJ/mol. Such a low value of $E_a$ indicates that the reorientational motions of the alkyl side group terminated with the thiol moiety are responsible for that. However, taking into account that the sum of the activation barrier for the conformation change of the alkyl chain ($E \sim 14$ kJ/mol) and the energy required to break S–H:S bonds ($E_{S-H:S} \sim 6$ kJ/mol) is comparable to the value of $E_a$, the $\gamma$-process in PMMS, one can assume that this secondary relaxation process is in some way indirectly connected to the dynamics of the hydrogen bonds.

The data reported herein clearly indicated that one has to be careful in the assignment of the molecular origin of the relaxation processes detected in dielectric data of the polymers tending to form supramolecular domains ordered on the nanoscale. Even a close correspondence between relaxation times obtained for a given relaxation process estimated from different techniques does not necessarily mean that they have the same molecular origin.

**CONCLUSION**

In this paper, we have discussed the results of complementary structural, mechanical, and dielectric studies performed for poly(mercaptopropyl)methylsiloxane (PMMS) characterized by chain backbone of alternate silicon and oxygen atoms substituted by a polar pendant group terminated with the thiol/sulfanyl (–SH) moiety. Interestingly, the performed FTIR and XRD measurements indicated that the examined PMMS forms weak but relatively efficient directional hydrogen bonds between neighboring chains, in a direction perpendicular to the chain backbone and, in consequence, supramolecular lamellar domains. Surprisingly, the collected dielectric spectra revealed the presence of the two relaxation processes at high temperatures, which merge upon cooling. Considering the outcome of XRD and rheological investigations, it was found that $\alpha'$ (the slower process of Debye-like shape) and $\alpha$ (the faster one) processes detected in loss spectra are connected with the association–dissociation process of the locally ordered nanolamellar domains connected via H bonds and segmental motions, respectively. However, it should be mentioned that the collected data do not entirely exclude the sub-Rouse origin of the observed $\alpha'$-process in PMMS. Data presented herein can be useful to understand the peculiar dynamical properties of associating polymers tending to form mesoscale self-assemblies.

**ASSOCIATED CONTENT**

Supporting Information

The Supporting Information is available free of charge at https://pubs.acs.org/doi/10.1021/acs.macromol.0c01289.

Experimental details and additional figures including correlation functions, FTIR spectra, the real part of dielectric spectra with the temperature dependences of $\tau$ and $\varepsilon'\varepsilon''\varepsilon''$, comparison of normalized shear and dielectric $\alpha$-peaks, experimental details and additional figures including correlation functions, FTIR spectra, the real part of dielectric spectra with the temperature dependences of $\tau$ and $\varepsilon'\varepsilon''\varepsilon''$, comparison of normalized shear and dielectric $\alpha$-peaks, dielectric loss spectra recorded for PMDS of $M_n \sim 4000$ g/mol, temperature dependences of $\tau$ determined for PDMS, PMMS, and HO-PMDMS-OH (PDF)

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Notes
The authors declare no competing financial interest.

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