CRYSTALLIZED STATE OF POLYMER IN ITS DISPERSION BEHAVIOUR

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INTRODUCTION

In the field of crystalline polymers the morphological and structural studies of polymers have remarkably developed since the fold structure of single crystal of polymers was discovered by Keller\(^1\) in 1957. This discovery has rendered us a new structural view, which is contributing to the interpretation of physical properties of crystallized polymers. Strict analysis of molecular arrangements within polymer crystal and beautiful electron photomicrographs of lamellar structure as observed in bulk crystallized linear polyethylene seem to admit no question of the state of aggregation of polymer molecules in the crystalline texture. This view is, however, not adequate in the case of polymers such as polyester, polyamide and polypropylene having 40–50 per cent amorphous region in their bulk-crystallized texture. Such polymers are rather more frequently found than highly crystalline polymers such as high density polyethylene and polyoxymethylene. For the polymers with medium degree of crystallinity the structural studies using x-ray diffraction, electron photomicrography and other methods seem to be insufficient to describe the state of aggregation of molecules in the crystalline texture, especially in the amorphous region. Dispersion behaviour of polymers gives information different from the structural studies in some point in grasping the actual state of crystallized polymer. The main purpose of this article is to describe the real state of crystallized polymer by supplementing the information of structural studies with knowledge obtained from dispersion behaviours.

AGGREGATION OF MOLECULES AND DISPERSION PHENOMENA

Molecular chains under less restraint in the crystalline texture initiate their thermal motions at a lower temperature or at a shorter time (higher frequency), while molecular chains under greater restraint initiate their thermal motion at a higher temperature or at a longer time. This means that the difference in the state of aggregation of molecules could be discussed by measuring the change of dispersion behaviour. Temperature dependence of relaxation time or activation energy for relaxation and shape of relaxation spectrum also allow us the more detailed discussion on the state of aggregation of molecules.

Relaxation mechanisms are detected by measuring dispersion or absorption phenomena. As for viscoelasticity the measurements of transient phenomena such as stress relaxation and creep are useful, while dispersion of storage modulus and absorption of loss modulus obtained by the dynamic
MOTOWO TAKAYANAGI

method using periodic excitation are frequently more sensitive and effective to achieve the present purpose. When the average relaxation time $\bar{\tau}$ is assumed to be characteristic of some type of molecular motion, the initiation of this motion can be sensitively detected by observing the dispersion of storage modulus $E'$ and the absorption of loss modulus $E''$, which appear under the condition that $\omega \bar{\tau} = 1$ is satisfied by changing frequency at a definite temperature or by changing temperature, that is, by changing $\bar{\tau}$ at a definite frequency (isochronal measurement).

As a typical example of dispersion curve of the crystalline polymer, that of poly-4-methyl-pentene-1 is cited in Figure 1\textsuperscript{28}. From the high temperature

![Figure 1. Tensile storage modulus $E'$ and loss modulus $E''$ vs. temperature at 110 c/s for three kinds of samples of isotactic poly-4-methyl-pentene-1 [(1) Bulk-crystallized sample quenched at $-70^\circ\text{C}$ from the melt, (2) Annealed sample at 210°C for 15 h, (3) The single crystal mat] side, the crystalline absorption is found at about 170°C, which increases its intensity with increasing crystallinity, the primary absorption associated with the initiation of the micro-brownian motion in the amorphous region is found at about 50°C, which decreases its intensity with increasing crystallinity. The secondary absorption associated with the local twisting motion of the main chains is latent in the background of the primary absorption. Generally speaking, the secondary absorption of vinyl compounds absent from any flexible side chain such as isotactic polypropylene, polystyrene and polyvinyl alcohol are similarly latent, appearing only as a broad weak shoulder of the primary absorption, while the secondary absorptions of high density polyethylene and polyoxymethylene without side chains are usually very remarkable. The secondary absorption associated with the thermal motion of side chain is found at about $-160^\circ\text{C}$ at 110 c/s distinctly for poly-4-methyl-pentene-1.

Crystalline texture will be roughly divided here into the crystalline and the amorphous region. Discussions will be given of each region referring to some recent works of the author's laboratory. In addition to these discussions, the mechanical coupling of the amorphous and the crystalline region will be taken up as another subject of this article. The change of dispersion be-
CRYSTALLIZED STATE OF POLYMER IN ITS DISPERSION BEHAVIOUR

haviours caused by a different way of coupling of the two regions is consi-

DERSON ASSOCIATED WITH CRYSTALLINE REGION

Structurally defined polymer crystals and their dispersion behaviours

As polymer single crystal has a high degree of crystallinity compared with

measurements of viscoelasticity of single crystal were made in the form of

Crystalline absorption of the single crystal mat is affected by the morpho-

logical factors of crystal and its perfection. Single crystal samples of linear

polyethylene with different thickness L were prepared by strictly regulat-

ing the conditions of isothermal crystallization. Lamellar thickness L was

evaluated by the long period of x-ray small angle scattering. Figure 2 shows

the systematic increase of intensity of the crystalline absorptions and the

shift of the dispersion temperature towards the high temperature side with

increasing lamellar thickness from 94 (curve 1) to 167 Å (curve 5). By

changing the frequency of measurements and applying the method of

reduced variables the master curve of $E''$ is obtained and from this curve the

relaxation spectrum can be evaluated with the usual approximate method. Figure 3 shows the relaxation time spectra of the single crystal mats with

lamellar thickness of 94, 104, 114, 137 and 167 Å. The average relaxation
time $\tau$ shifts to the side of longer relaxation time with increasing lamellar

thickness. This tendency will be qualitatively understood by considering

the corresponding modes of torsional motion of molecular segments con-

stituting the different lamellar thickness. The more elaborated theory has

been developed by Okano. According to Okano's theory the tendencies of

the increased intensity of relaxation and the increased absorption tempera-
ture with increasing thickness are expressed by the linear relationships of

$\log L$ vs. reciprocal temperature of the crystalline absorption and of the

average relaxation time $\tau$ vs. $L^2$. However, definite discrepancies were found

between our experimental results and the relationships predicted by Okano.

The shape of the relaxation spectra of the crystalline absorption is very broad

when compared with that of the primary absorption of the completely

amorphous polymer. The interpretation of such broad spectra by the

molecular theory is a problem to be solved in the future, while the broadness

is considered to be caused by the coupled oscillation of many molecular

chain segments composing a crystal block in single crystal.
It is well known that the single crystal of polyethylene thickens by heat-treatment above its crystallization temperature. Figure 4 shows absorption curves for the annealed single crystal mats of polyethylene. With increasing lamellar thickness determined by long period of small angle x-ray diffraction, the absorptions of the samples shift to the high temperature side in the same way as in the case of the isothermally crystallized samples. It should be noticed, however, that the absorption intensity of the crystalline absorption of the annealed samples decreases with increasing thickness contrary to the case of the isothermally crystallized samples. Figure 5 shows a plot of the absorption temperature $T_{max}$ against long period $L$. The relation is very similar to the relation of melting temperature of polyethylene crystal evaluated by Lauritzen and Hoffman or of $n$-paraffin evaluated by Broadhurst to $L$. The $T_{max}$ of both the samples obtained by isothermal crystallization and by annealing is expressed by a single curve as a function of long period. On the other hand, the $E''_{max}$ values of the isothermally...
Figure 3. Relaxation spectra at 60°C for the single crystal mats with long periods of 94, 104, 114, 137 and 167 Å which are prepared from the composite curve of $E''(\omega)$ based on the measurements at different frequencies and temperatures.

Figure 4. $E''$ vs. temperature at 110 c/s for the single crystal mats of linear polyethylene annealed at 70°, 90°, 100°, 115° and 128°C, corresponding to the long periods of 102, 127, 156, 247 and 473 Å. Original single crystal prepared at 40.5°C is 101 Å thick.
Figure 5. Temperature of the crystalline absorption ($\alpha_c$) maximum vs. long period for the isothermally crystallized polyethylene single crystal (●) and the annealed samples starting from the single crystals A and B (○, ⊗). Melting temperature $T_m$ vs. long period (— — —) for polyethylene single crystal after Lauritzen and Hoffman's equation. The crystalline absorption ($\alpha_c$) temperature associated with the irreversible structural change$^{15}$ vs. long period for the isothermally crystallized samples (○).

The crystallized samples and the annealed ones are not expressed by a single curve but by two separate curves of quite different types as shown in Figure 6. The only possible interpretation for these contradictory relationships of $T_{max}$ ($\alpha_c$) and $E''_{max}$ ($\alpha_c$) against long period $L$ is that the value of $T_{max}$ is primarily determined by the thickness of crystal with almost perfect

Figure 6. Maximum intensity of the crystalline absorption $E''_{max}$ vs. long period for the isothermally crystallized samples (○) and the annealed samples (●, ○). Results of two series of annealed samples are shown for the original crystals A and B.
CRYSTALLIZED STATE OF POLYMER IN ITS DISPERSION BEHAVIOUR

regularity, but, on the other hand, the absorption intensity \( E''_{\text{max}} \) is determined by the quantity of the crystal of the thickness \( L \). If so, the quantity of such a perfect crystal region with thickness \( L \) should be decreased by increased annealing temperature.

The support of this view from the structural or morphological studies is not enough to confirm the view. There are, however, some experimental facts concerned with it. According to Holland\(^9\) the moife pattern of the annealed polyethylene single crystal has a very much distorted aspect, indicating that the concentration of dislocation is increased by a factor of one thousand as compared with that of the original crystal. In such a case the crystal part with perfect regularity will be largely decreased. The \((110)\) plane used in their observation is considered to be distorted like an amoeba shape and the area showing the regular stripes as observed in the isothermally crystallized samples evidently decreased to a fraction of the original ones. Such a structural change will be enough to interpret the decrease of the absorption intensity with increasing \( L \) as shown in Figure 6. The nature of the defect region is also discussed by Kedzie\(^10\), who has found the fact that the narrow component of the differential curve of n.m.r. absorption of the single crystal shows a remarkable anisotropy, i.e. the second moment takes a maximum value when the magnetic field direction accords with that of molecular axis in the crystal. The broad component has two maxima depending on the orientation of the crystal. The fact that the narrow component has anisotropy means that the defect region shows preferable orientation along the direction of molecules in the crystal and that it exists within the crystal phase. We assume here that the defect regions are included in the lamellae thickened by heat-treatment and they do not contribute to the crystalline absorption. Slichter\(^11\) found that the narrow component appears by heat-treatment of single crystal of polyethylene and it increases its intensity with increasing rate by raising the temperature of annealing. This fact corresponds well with the observation that the \( E''_{\text{max}} \) of the crystalline absorption decreases with increasing annealing temperature.

It is known that the broad component of n.m.r. spectrum of the solution grown polyethylene crystal narrows at a certain temperature range with increasing temperature. According to Iwayanagi\(^12\) the temperature at which the broad component begins to narrow rises with increasing crystallization temperature. This means that the molecular motions in the crystal region of perfect regularity tend to initiate at the lower temperature for the thin crystal obtained at the low crystallization temperature. This tendency is in accordance with our observation that the \( T_{\text{max}} \) shifts to the high temperature side with increasing crystallization temperature as shown in Figure 4. Sinnott\(^13\) has recently reported on the mechanical relaxations in single crystal mat of polyethylene; his results are almost the same as ours in that the crystalline absorption of the annealed and thickened crystal shows the decreased relaxation intensity (expressed by Sinnott with the absorption area instead of with the absorption maximum reported by us). Based on this observation Sinnott proposed that the crystalline absorption is associated with the folded region attached to the surface of the thickened lamella. This interpretation seems to encounter the difficulties in interpreting our
observation that the crystalline absorption intensity increases with increasing long period for the isothermally crystallized samples with different thicknesses. Mechanical dispersions associated with the loose loops attached to the crystal surface will be discussed in more detail under the next section, discriminating from the crystalline absorption mentioned here.

In the above discussion the maximum intensity of the absorption $E''_{\text{max}}$ was employed for convenience as the measure of the magnitude of an absorption. The magnitude of dispersion $\Delta E$ is given by the following equation based on the interrelation of viscoelastic functions.

$$\Delta E = \int_{-\infty}^{\infty} H_1 \, d \ln \tau = \frac{2}{\pi} \int_{-\infty}^{\infty} E''(\omega) \, d \ln \omega.$$ 

Evaluation of $\Delta E$ values of the isothermally crystallized samples was made using the relaxation spectra of Figure 3 and compared with the $E''_{\text{max}}$ value of Figure 2. Satisfactory proportionality was obtained between both values of $E''_{\text{max}}$ and $\Delta E$ corrected for the recrystallization effect.

It should be noticed, however, that the absorption curves of the annealed single crystal and the bulk crystallized sample$^{2,15}$ could not be reduced to prepare the composite curves of $E''$ due to the appearance of the new crystalline absorption ($\alpha_{\text{c}}'$) in addition to the $\alpha_{\text{c}}$ absorption at its low temperature side$^{15}$. We did not try here to evaluate these two absorptions separately. $E''_{\text{max}}$ value was employed as the measure of magnitude of an absorption in the above discussion.

Next we would like to proceed to the subject of crystalline texture of the bulk crystallized samples apart from the problem of single crystal. It is well known that the spherulite is composed of piled thin plate-like crystals (lamella.), which are very similar in their structure to the single crystal except an existence of tie molecules between neighbouring lamellae in the spherulite. The foregoing discussions on the single crystal will be, therefore, reasonably applied to the crystalline texture of the bulk crystallized sample under some restrictions.

Broken lines of Figure 2 are the $E'$ and $E''$ vs. temperature curves at 110 c/s for the bulk crystallized polyethylene. Compared with the isothermally crystallized single crystal, the bulk crystallized sample rather resembles the annealed single crystal in the shape of its dispersion curves. According to our concept of the crystallized state of isothermally crystallized single crystal and the annealed one mentioned above, the lamellar phase of spherulite includes a somewhat large defect region, by which the intensity of the crystalline absorption is largely decreased. And at the same time the low temperature absorption at about $-120^\circ\text{C}$ of the bulk crystallized sample is remarkably increased compared with that of the single crystal samples as if it compensates the decreased magnitude of the crystalline absorption. The $-120^\circ\text{C}$ absorption is called here the $\beta_{\text{a}}$ absorption, which is closely related to the amounts of defect region in the crystal and of the frozen amorphous region. The same relation can be found in the absorption curves of polyoxymethylene between the samples of the single crystal mat and the bulk crystallized film$^{16}$. 

562
CRYSTALLIZED STATE OF POLYMER IN ITS DISPERSION BEHAVIOUR

The authors have recently found an experimental support for the concept that the low temperature secondary absorption associated with side chain motion $\beta_{sc}$ (some authors call it $\gamma$) is divided into two components; the one is concerned with the frozen amorphous region ($\beta$-A) and the other with the defect region in the crystalline phase ($\beta$-C). The secondary absorption at about $-155^\circ$C of the bulk crystallized isotactic poly-4-methylpentene-1 (P4MP1) splits into two separable regions; the higher temperature one is concerned with the frozen amorphous region, its temperature location is in accordance with that of the completely amorphous atactic samples and its magnitude (the area under the $E''$ vs. $1/T$ curve) decreases in proportion to the fraction of the amorphous region or with increasing crystallinity ($\chi$), and the lower temperature one is concerned with the defect region of the crystalline phase, although its magnitude (the area of the absorption separated from that of the frozen amorphous region) is not necessarily proportional to the degree of crystallinity $\chi$, but amounts to some fraction of $\chi$ depending upon the perfectness of crystalline region. It should be noted that the secondary absorption of P4MP1 at about $-155^\circ$C is concerned with the thermal motion of side chain ($\beta_{sc}$) and not with the local twisting motion of main chain ($\beta_a$) as mentioned in the introduction. The separation of the $\beta_{sc}$ absorption of P4MP1 into the one associated with the frozen amorphous region $\beta_{sc}$-A and the defect region $\beta_{sc}$-C is illustrated in Figure 7. The fraction of the amorphous region ($A$) can be evaluated as the ratio of the area of the $\beta_{sc}$-A absorption to that of the atactic sample. Defect region ($A'$) can be evaluated by the value of $A$ multiplied by the ratio of the magnitude of $\beta_{sc}$-C absorption to that of the $\beta_{sc}$-A absorption. For example the bulk crystallized P4MP1 with degree of crystallinity $\chi = 0.46$ evaluated by the x-ray method gives the values of $A = 0.56$ by the absorption method, the absorption ratio of $\beta_{sc}$-C/$\beta_{sc}$-A = 0.28/0.72, and therefore $A' = 0.56 \times 0.28/0.72 = 0.22$. The fraction of perfect crystal C is given by $1 - A - A' = 0.22$. The value of $A'/(1 - A)$, the fraction of the defect region in lamellar phase, is evaluated as 0.50, i.e. the defect region amounts to about a half of lamella. The $A'/(1 - A)$ value depends upon the crystallization condition of the sample, and its low value corresponds to a high degree of perfection of crystal regularity. The case of P4MP1 is rather fortuitous, as its secondary absorption is separable into two regions. Usually two such regions are united into one secondary absorption.

Another evaluation method of defect region in the lamellar phase of the bulk crystallized sample was previously presented by the authors in the case when the primary absorption (called here $\alpha_a$) associated with the initiation of micro-Brownian motion of the main chains belonging to the amorphous region can be found. This method is also based on the same concept of the secondary absorption as mentioned above. There are two kinds of secondary absorption; $\beta_{sc}$ and $\beta_a$. The latter is concerned with the local twisting motion of main chains and is often remarkable for the polymer absent from side chains. The treatment of $\beta_a$ is the same as that of $\beta_{sc}$.

The crystalline texture of the bulk crystallized polymer is divided again into three regions, A, A' and C, corresponding to the degree of thermal motion of molecules. The region A is defined as the region, the molecules in which are capable of the micro-Brownian motion above its glass transition
temperature. The region $A'$ is defined as the region, the molecules in which are only capable of the local twisting motion but incapable of diffusional motion even if above $T_g$. Such circumstances will be realized only within the lamellar phase. The region $C$ is the crystalline region with perfect

\[
\begin{align*}
A &= \frac{(\Delta E)_\alpha}{(\Delta E)_\alpha} \\
A' &= \frac{(\Delta E)_\beta}{(\Delta E)_\beta} - A \\
C &= 1 - A - A'
\end{align*}
\]

relationships (1), (2) and (3), where $\Delta E$ is the magnitude of dispersion, and the denominator represents the quantity referring to the wholly amorphous sample and is evaluated by extrapolating the absorption magnitude to

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**Figure 7.** $E''$ vs. $1/T$ at 110 c/s for the crystallized isotactic poly-4-methyl-pentene-1 (upper curve) and the atactic one in a temperature range of low temperature secondary absorption $\beta_{sc}$. The secondary absorption $\beta_{sc}$ is divided into two regions $\beta_{sc-A}$ (high temperature absorption) and $\beta_{sc-C}$ (low temperature one).

The right side is the diagram representing the relationship among the degree of crystallinity $\chi$, the region $A$ evaluated with the $\beta_{sc-A}$ absorption, the region $A'$ with the $\beta_{sc-C}$ absorption, and the perfect crystal region $C = 1 - A - A'$. $A$ is the fraction of the frozen amorphous region and $A'$ is the defect region in lamella. Confer the evaluation method in the text.

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564
CRYSTALLIZED STATE OF POLYMER IN ITS DISPERSION BEHAVIOUR

the zero crystallinity. **Figure 8** schematically represents the relation among degree of crystallinity \( x \) determined by physical methods such as density or x-ray, and the viscoelastically defined regions, \( A, A' \) and \( C \).

Polyethylene terephthalate is a polymer suitable for application of the above method, since this polymer displays a remarkable \( \alpha_a \) absorption and a \( \beta_a \) absorption at the same time. The author\(^{18} \) employed the \( \alpha_a \) absorption maximum \( E'' \max \) instead of \( \Delta E_a \) in Eq. (1). Illers\(^{20} \) has pointed out that the absorption magnitude should be evaluated by the area under the \( E'' \) vs. \( 1/T \) curve. More detailed inspection of the \( \alpha_a \) absorption has been made by the author\(^{18} \) based on the relaxation spectra for the samples with different crystallinity. The shape of the relaxation spectra in a primary absorption region of the crystallized sample is not simple and suggests some composite relaxation mechanisms. At present the \( \alpha_a \) absorption magnitude can be more strictly evaluated in the dielectric absorption by using the area under the \( e'' \) vs. \( \log f \) curve. The magnitudes of \( \langle \Delta E \rangle_a \) and \( \langle \Delta E \rangle \beta \) were plotted against degree of crystallinity \( x \) for various kinds of polyesters\(^{21}, \, 22 \). The \( \alpha_a \) absorption decreases in magnitude remarkably with increasing \( x \), while the magnitude of \( \beta_a \) absorption does not decrease so much even when extrapolated at \( x = 1 \). **Figure 8** shows the schematic representation of the

![Diagram](attachment:diagram.png)

**Figure 8.** Schematic representation of the relationship among degree of crystallinity \( x \), the viscoelastically evaluated regions, \( A, A' \) and \( C \) for the bulk crystallized polyethylene terephthalate with \( x = 0.60 \). For evaluating the regions \( A, A' \) and \( C \) the magnitudes of the primary \( (\alpha_a) \) and the secondary \( (\beta_a) \) absorption were used. Related domains of these absorptions are indicated in the figure.

fractions of \( A, A' \), and \( C \) regions of polyethylene terephthalate with \( x = 0.60 \). Referring to these evaluations, an assignment of the regions \( A, A' \) and \( C \) to the real structure of spherulite will be made as shown in **Figure 9**. The region \( A' \) is assigned to the defect region within the lamella and the region \( A \) is assigned to the amorphous region segregated from the lamellar phase. The model of the lamellar structure, which includes a large defect region depending on circumstances amounting to about a half of the lamellar phase, does not seem to be in conflict with our concept on the structure of the annealed and thickened single crystal.

At last the effect of the chemical structure of polymers upon the crystalline absorption will be discussed.\(^3 \) Single crystal mats of various polymers such as poly-\( \alpha \)-olefins, polyethers and polyvinylalcohol give the remarkable crystalline absorptions on the contrary to the bulk crystallized samples prepared from the same polymers. **Figure 10** shows the temperature dependence of \( \tan \delta \) at 138 c/s for the two kinds of solution-grown samples and the bulk-crystallized sample of isotactic polybutene-1\(^{23} \). In the single crystal
samples the primary absorption at about 0°C is remarkably depressed to be
detectable only as a shoulder and the prominent crystalline absorption
newly appears at 80° or 100°C depending on the conditions of crystallization.

Activation energies $\Delta H^*$ evaluated by the temperature dependence of the
crystalline peak position at different frequencies for several polymers,
activation volumes $\Delta V^*$ and their melting temperatures $T_m$ are listed in

Figure 9. Model for the local structure of spherulite. The amorphous region A is the region
segregated from the lamellar phase, the region $A'$ is the defect region in the lamellar phase
and the region C is the lamellar phase with perfect crystal regularity.

Table I. It should be noted that the value of PVA is considered to correspond
to the unstable crystal form like smectic or nematic structure. The solution-
cast film was used for PVA sample.

The fact that the activation energy for relaxation of the crystalline region
increases with increasing melting temperature leads us to the conclusion
that the factors affecting the molecular motion associated with the crystalline
region are common to those factors determining the thermodynamical
stability of the corresponding crystals.

The fact that the typical crystalline absorption is closely related to the
melting temperature concerned with the chemical structure and at the same
time with the morphological factor strongly suggests that the crystalline
absorption is associated with mechanical relaxation of the crystalline region
with high regularity.

**Structurally undefined low order crystals and their dispersion
behaviours**

There is a possibility of existence of a phase with intermediate regularity
between the well-defined crystalline region and the amorphous region. Such
a phase is usually difficult to describe crystallographically.

*Figure 11* shows the tan $\delta$ vs. temperature curves of the four kinds of
samples of Nylon 6. In the solution grown crystal of Nylon 6 there is in its
temperature dispersion curve a broad absorption at about 120°C at 110 c/s
in addition to the typical crystalline absorption found at about 200°C.
CRYSTALLIZED STATE OF POLYMER IN ITS DISPERSION BEHAVIOUR

Figure 10. Tan δ vs. temperature at 138 c/s for the bulk crystallized sample (○) and the solution grown samples (● and ●) for isotactic polybutene-1

Table 1. Activation energies (ΔH*), activation volumes (ΔV*), and number of chain atoms (N) concerned with crystalline relaxation

<table>
<thead>
<tr>
<th>Polymer</th>
<th>ΔH* (kcal/mole)</th>
<th>Tc (°C)</th>
<th>Tm (°C)</th>
<th>ΔV* (ml/mole)</th>
<th>V (ml/mole)</th>
<th>$\frac{N = \Delta V^*}{V}$</th>
</tr>
</thead>
<tbody>
<tr>
<td>PB-I</td>
<td>25</td>
<td>70</td>
<td>97°c</td>
<td>520</td>
<td>29-4°d</td>
<td>18</td>
</tr>
<tr>
<td>PE</td>
<td>30</td>
<td>90</td>
<td>135</td>
<td>550</td>
<td>13-9</td>
<td>40</td>
</tr>
<tr>
<td>PVA</td>
<td>48</td>
<td>130</td>
<td>—</td>
<td>800</td>
<td>16-4</td>
<td>49</td>
</tr>
<tr>
<td>PP</td>
<td>48</td>
<td>120</td>
<td>165</td>
<td>820</td>
<td>22-4</td>
<td>37</td>
</tr>
<tr>
<td>POM</td>
<td>55</td>
<td>160</td>
<td>180</td>
<td>860</td>
<td>9-9</td>
<td>87</td>
</tr>
</tbody>
</table>

a ΔH* is evaluated from the change of absorption maximum with frequency.
b ΔH* is evaluated from the temperature dependence of shifting factor aT.
c Melting temperature of modification III of PB-I determined by DTA method.
d V is evaluated by the lattice constants of Mod III obtained by the authors; tetragonal, $\alpha = b = 14.88\text{Å}$, $c = 12.31\text{Å}$, $7_1$ helix.
The 200°C absorption cannot be usually detected for the bulk crystallized samples of Nylon 6. In the bulk crystallized samples the remarkable primary absorption (the \( \alpha_a \) absorption) appears at about 80°C instead of the broad 120°C absorption of the solution-grown crystal. According to our view, the 120°C absorption of the bulk crystallized sample is considered to be still latent and unseparable, being in the high temperature side of the \( \alpha_a \) absorption. This tendency can be more clearly seen in the \( \tan \delta \) curve of the bulk crystallized sample in Figure 11 prepared by melting it at 300°C, crystallizing it slowly at a high temperature close to the melting temperature, and removing about 19 per cent of produced \( \varepsilon \)-caprolactam by hot water. In this sample it can be shown that the 120°C peak coexists with the 80°C absorption.

In even Nylons such as Nylon 6, 8 and 12\(^{25} \), when they are treated by iodine–potassium iodide solution and the iodine absorbed is rapidly removed by treating with alcoholic sodium thiosulphate solution, the dispersion curve similar to that of the solution grown sample is obtained, i.e. the \( \alpha_a \) absorption at about 80°C is largely depressed and the 120°C absorption remarkably increased as seen in the open circle curve in Figure 11. In this case iodine acts at first to break the hydrogen bonds in crystal and rapid desorption of iodine rebuilds the hydrogen bonds in a somewhat random way. This process seems to be closely related to the appearance of the 120°C peak for Nylon 6. The behaviours of Nylon 8 and 12 are very similar to those of Nylon 6.

On the other hand the odd Nylons such as Nylon 7, 9 and 11 give only the primary absorption at about 70°C without any background portion at its high temperature side even if the iodization and desorption treatment have been carried out\(^{25} \). Figure 12 shows the \( \tan \delta \) vs. temperature curve of Nylon 9. It should be mentioned that any structural evidence using x-ray diffraction or infrared absorption cannot be found at present for the existence of the phase characteristic of the 120°C absorption. It is regrettable that the
authors are still at the stage of speculation for interpreting these problems. One possible interpretation of the existence or absence of the 120°C absorption in even and odd Nylons will be presented next.

![Graph showing tan δ vs. temperature for three kinds of samples of Nylon 9](image)

**Figure 12.** Tan δ vs. temperature at 110 c/s for three kinds of samples of Nylon 9 [(●) bulk crystallized sample annealed at 180°C, (○) iodine-adsorbed and desorbed sample, and (◆) is for the same sample annealed at 160°C after iodine is desorbed]

Even Nylon has a special crystal form or γ, which is composed of shortened fibre period to saturate all the hydrogen bonds between neighbouring parallel-directioned molecules, in addition to the usual crystal forms of α and β, which are composed of planar zigzag chains to saturate the hydrogen bonds between the anti-parallel directioned molecules. On the other hand odd Nylons have no necessity of assuming the γ form in addition to the normal α and β crystal forms, since both parallel-directioned and anti-parallel-directioned molecules can completely saturate their hydrogen bonds between neighbouring molecules in an extended planar zigzag form.

From such structural factors, if iodine is removed rapidly from the iodized samples, there is a possibility that for even Nylons both parallel and anti-parallel-directioned molecules will form hydrogen-bonds in a mixed state, where the different fibre periods disturb the formation of the well defined crystal modification such as α or γ form and as its consequence a phase of low regularity such as nematic structure will be formed. On the other hand for odd Nylons there will be no possibility of formation of nematic structure. The nematic form will behave almost in the same way as in the so-called amorphous region, while somewhat higher regularity of the nematic region will be enough to shift its absorption peak towards the high temperature side of the primary absorption.

It is worth mentioning that Nylon 66 has a possibility of formation of nematic structure in spite of the absence of γ modification. In the case of...
Nylon 66 there are two ways of arrangement of neighbouring planar zigzag molecules in the hydrogen bonded sheet: one way is to saturate all the hydrogen bonds between neighbouring molecules, and another way is realized by shifting the one planar zigzag chain along the neighbouring one by an amount of one residual radical, as the consequence of which only one half of the hydrogen bonds in the former case is saturated. Figure 13 shows two kinds of arrangements of molecules in Nylon 66. These two molecular arrangements will be realized in a mixed state to form the nematic structure when the regeneration from the iodized state is rapidly processed. Figure 14 shows that the iodine-desorbed Nylon 66 has an increased absorption at about 150°C, while the crystallized sample from the melt gives only a large primary absorption at about 60°C as is well known.

Polyvinyl alcohol (PVA) has an absorption at about 120°C, which locates between the primary absorption at about 80°C and the crystalline absorption at about 200°C. These absorptions are found in the solution-cast film. The crystalline absorption is remarkable especially for the annealed solution-grown crystal. The 120°C absorption increases its intensity for the samples partially acetylated or partially formalized as shown in Figure 15. Such treatments will increase the nematic region in the crystalline texture of PVA.

Natta reported the smectic modification of isotactic polypropylene (PP) of pseudo-hexagonal form. The authors have found in the dispersion curve of the quenched samples of PP the absorption characteristic of smectic
structure at about 50°C, being distinctly separated from the primary absorption locating at 20°C at 110 c/s. Figure 16 shows the effect of treatment on the appearance of the 50°C absorption characteristic of smectic modification. Dilatometry proves that the melting temperature of the smectic structure is 50°C. The process of remelting the smectic structure just above 50°C was reflected in an increase of the primary absorption, and the process of recrystallization by heat-treatment of the remelted sample was reflected in disappearance of the 50°C absorption being accompanied by a decrease of the primary absorption at 20°C together with an increase of crystallinity. Structural change of these processes studied by x-ray diffraction well accorded with the change of absorption curves. The crystalline absorption of PP can be found at about 120°C, especially distinctly for the solution-grown crystals27.

The absorption associated with the unstable modifications are also observed for the solution grown crystals of poly-4-methyl-pentene-1 at about 120°C as shown in Figure 1728, for the solution grown crystals of poly-n-pentene-1 at about 50°C29, for the single crystal of poly-3,3-bischloromethyl oxacyclobutane at about 60°C while its a absorption is found at 30°C30.

571
The crystal structure characteristic of these absorptions could not be found out crystallographically, while the regions characteristic of these absorptions are clearly different from those associated with the primary absorption. Such a structurally undefined but somewhat more ordered region than the amorphous region is to be taken into account to understand the actual crystallized state of polymers.

**DISPERSION ASSOCIATED WITH AMORPHOUS REGION**

The reasoning that the state of the amorphous region of the bulk crystallized polymer is different from that of the completely amorphous polymer will be partly supported by the fact that the relaxation spectrum associated with the primary absorption is always much broader than that of the amorphous polymer. Typical example is found in quenched and annealed polyethylene terephthalate, the degree of crystallinity of which can be regulated from 5 to 45 per cent\(^{18}\). The broadness of the spectrum is considered to be caused by the circumstances that the molecules within the amorphous region of the crystallized polymer suffer from various degrees of restraint by the surrounding lamellae.

The reasoning that the molecules causing the primary absorption are in a state of being segregated from the lamellar phase will be supported by the
CRYSTALLIZED STATE OF POLYMER IN ITS DISPERSION BEHAVIOUR

The fact that the primary absorption of the crystallized isotactic polypropylene (PP) locates at almost the same temperature as that of the isotactic PP blended with atactic PP\textsuperscript{8,16}. Atactic PP in the crystalline texture of the blend cannot be structurally incorporated into lamellae mainly composed of isotactic PP. The fact that the primary absorptions of both the samples resemble each other except their shapes means that the amorphous region of isotactic PP is not so much different from the state of atactic PP segregated from the lamellae in the blend system. Quantitative analysis based on this assumption has been given in the previous papers\textsuperscript{8,16}.

![Figure 16. Tan δ vs. temperature at 110 c/s for the quenched and the annealed isotactic polypropylene samples. The primary absorption is found at about 20°C and the absorption concerned with smectic structure is at about 50°C.](image)

Theoretical considerations on the primary dispersion mechanisms seem to support the foregoing concept of the amorphous region. The primary absorption is related to the initiation of the micro-Brownian motion of molecules belonging to the amorphous region, and referring to the molecular theory on the viscoelasticity of the completely amorphous polymer presented by Rouse and Bueche\textsuperscript{31}, some motional modes associated with the cooperative motion of fairly large length of molecules, which will not be

573
included within the lamellar phase, should be considered at the long time side or the high temperature side of the transition region of the crystalline polymer.

![Heat treatment of P 4MP1](image)

Figure 17. Tan δ vs. temperature at 110 c/s for the solution grown crystal mat of poly-4-methylpentene-1 and the samples annealed of it. The primary absorption (δ) is found at about 50°C. The 120°C peak is related to the unstable crystal form.

For the polymers being almost absent from the primary absorption such as linear polyethylene or polyoxymethylene, the prediction that the amorphous region accords with the defect region within the lamellar phase may be allowed. However, for the polymers displaying the definite primary absorptions such as polyamide, polyester and poly-α-olefins, it should be considered that the amorphous region associated with the primary absorption is segregated from the lamellar phase. In this view the single crystal of linear polyethylene prepared from xylene solution, in which the primary absorption is absent\(^2\)\(^3\), will be composed of the molecules, being sharply folded at the crystal surface to form tight loops, which are incapable of micro-Brownian motion.

At this stage it will be necessary to present an example of single crystal displaying the primary absorption, which suggests the possibility of formation of loose loops attached to the crystal surface. Single crystal of trans-1,4-polybutadiene (PBD) can be prepared from various solutions\(^8\). Figure 18 (given in the section printed on art paper) shows the electron photomicro-
CRYSTALLIZED STATE OF POLYMER IN ITS DISPERSION BEHAVIOUR

graph of single crystal of trans-1,4-polybutadiene prepared from benzene solution. Figure 19 shows that the dispersion curve of the mat of PBD single crystals (curve 2) has a remarkable primary absorption at about $-20^\circ$C at 110 c/s, being largely different from the single crystal mats of polyethylene and polyoxymethylene. By annealing the mat of PBD crystals at 100°C, the $-20^\circ$C primary absorption has disappeared completely and the tensile storage modulus has been increased (curve 3). Dispersion curves expressed by the broken lines of Figure 19 are for the bulk crystallized samples (curve 1). Figure 20 shows the results of detailed inspection of the effect of annealing temperature on the primary absorption of PBD single crystal. In the results of Figure 20 the loose loops associated with the primary absorption disappear suddenly by annealing at 57°C just above the crystal transformation temperature $55^\circ$C. The transformation temperature of the bulk crystallized PBD is known as $76^\circ$C. The decrease of the crystal transformation temperature from $70^\circ$C to $55^\circ$C observed here is entirely caused by the morphological difference of lamella between the single crystal and the bulk crystallized sample. X-ray analysis proves that the crystal form of the single crystal sample changes from Form 1 to Form 2 at $55^\circ$C in the same way as the bulk crystallized one changes at $75^\circ$C. At the crystal transformation temperature $55^\circ$C the lamellar thickness determined by the area method jumps from 100 Å to 160 Å. Figure 21 shows the change of the crystal thickness evaluated by the line profile method applied to the meridional x-ray diffraction spot. The degree of crystallinity evaluated by the n.m.r. absorption method changes from 76 per cent of the single crystal mat to 91 per cent of the same sample annealed at 100°C. The narrow component of n.m.r. differential curve is largely decreased by heat treatment. All these facts

Figure 19. $E'$ and $E''$ vs. temperature for the three kinds of samples of trans-1,4-polybutadiene at 110 c/s [(---) is for the bulk crystallized sample, (O) is for the mat of single crystal, and (●) is for the single crystal mat annealed at 100°C]
Figure 20. Effect of annealing temperature on the tan δ vs. temperature curve at 110 c/s [Curve 1 is for the original single crystal mat of trans-1,4-polybutadiene, curves 2, 3, 4 and 5 are those for the single crystal mats annealed at 40°, 53°, 54°, and 57°C respectively. The crystal transformation temperature is 55°C]

Figure 21. Thickness of crystal phase of single crystal of trans-1,4-polybutadiene vs. annealing temperature. Thickness is determined by the line profile method applied to the meridional x-ray diffraction spot.

prove that the loose loops causing the primary absorption are attached directly to the single crystal surface and by heat treatment these loose loops are tightened, being accompanied at the same time by the thickening of the single crystal. Figure 22 schematically represents the models interpreting the phenomena of disappearance of the primary absorption and of thickening of lamellae. In Figure 22 the lamellar thickness was evaluated by comparison of the area of the single crystal domain before and after heat treat-
ment in the electron photomicrographs. The thickness of crystalline region of the original single crystal 72 Å was evaluated by the line profile of the x-ray diffraction spot near the meridian corrected for the instrumental broadening. The crystallinity of the original sample evaluated by the values of the total and the crystal portion thickness is 74 per cent after correction of density and accords well with the value of 76 per cent evaluated by n.m.r. absorption.

It is interesting that the average length of loose loops in the original crystal is evaluated from the above data as about 28 Å or 22 carbon atoms and this length is considered to be enough to initiate the micro-Brownian motion.

In order to survey the effect of the solvent power on the primary absorption of PBD single crystal it will be meaningful to discuss the mechanism of loose loop formation. Figure 23 shows the tan δ curves of single crystal mats prepared from benzene, cyclohexane, n-heptane and methyl isobutyl ketone (MIBK) at the crystallization temperatures of 18°, 24°, 62° and 75.5°C respectively. The intensity of the primary absorption in Figure 23 is increased in the order of MIBK, n-heptane, cyclohexane and benzene, i.e. with increasing solvent power. The solvent power was judged by the magnitude of the melting point depression according to Flory's equation. It predicts that the melting point depression by the solvent is determined by the solvent–polymer interaction parameter if the molar volume of the solvent is not so much different among the solvents used. A good solvent largely depresses the melting temperature as compared with a poor solvent.

A possible interpretation of the above results is that the free energy of formation of the critical nucleus for growth will be decreased, if the loops interact more with good solvents than with poor solvents to make loops loosen. Poor solvent has, on the other hand, a tendency of reducing the contact with the polymer segments to form rather tightened loops. Figure 24 shows a schematic representation of hypothetical growth mechanism with loose and tight loops in good and poor solvents respectively.

It is known that the primary absorption of bulk crystallized polymers does not disappear by heat-treatment. The primary absorption of bulk crystallized trans-1,4-polybutadiene represented by the broken lines in Figure 19 did not
disappear by heat-treatment at 100°C, while the primary absorption of the single crystal mat of the same polymer easily disappeared by the same treatment. This observation suggests that the existence of tie molecules between neighbouring lamellae largely resists the change of the state of the amorphous region of the bulk crystallized samples.

![Figure 23](image)

**Figure 23.** Effect of the solvents used to prepare the trans-1,4-polybutadiene single crystal upon the tan δ vs. temperature curve at 110 c/s. Benzene (○), cyclohexane (●), n-heptane (△), and methyl isobutylketone (●) are used at the crystallization temperatures, 18°, 24°, 62°, and 73.5°C respectively. Notice the tan δ peak at about —20°C corresponding to the primary absorption.

![Figure 24](image)

**Figure 24.** Schematic representation of hypothetical growth mechanisms with loose and tight loops in good and poor solvents.

The model of spherulite in its detailed structure presented by the authors has already been shown in Figure 9. Palmer and coworkers have first shown that the polyethylene spherulite gives well-defined crystal plates like single crystal after etching with fuming nitric acid. A similar experiment
CRystallized state of polymer in its dispersion behaviour

has recently been tried on the bulk crystallized sample of isotactic polypropylene by Minami, who treated this sample with fuming nitric acid and measured the viscoelastic properties of the etched residues embedded in the cellulose acetate. Figure 25 shows that the remarkable primary absorption

![Image](image_url)

Figure 25. Effect of etching with fuming nitric acid on bulk crystallized isotactic polypropylene (Minami) [(•) is for the sample untreated, (○) is for the sample etched by nitric acid, and (---) is for the mat of solution grown crystals prepared from xylene]

d of the original sample almost disappeared and the small 120°C crystalline peak of the original sample was overwhelmingly increased after nitric acid treatment. It is interesting that the absorption curve of the etched sample shows almost the same shape as that of the solution-grown crystal of polypropylene.

The crystallized state of polymer will be clarified in more detail if a knowledge of molecular motion from dispersion behaviour is added to the information from the structural and morphological studies.

**Dispersion curves and higher order structure**

Higher order structure of the bulk crystallized polymer is considered to be composed of the spherulitic lamellae and the amorphous region filling interlamellar space. Above the glass transition temperature the bulk crystallized sample resembles in some point high impact plastics, which are
composed of rubber particles dispersed in the matrix of hard plastics. The high impact interpolymer resin shows the rubber absorption at the low temperature and the plastics absorption at the high temperature of its dispersion curve. In the bulk crystallized polymer the primary absorption and the crystalline absorption can be observed in its dispersion curve, as if the amorphous region and the crystalline region correspond to the rubber and to the plastics phase of the interpolymer respectively. The way of mechanical connection of the amorphous and the crystalline region can be revealed by a detailed inspection of the dispersion curve. Such a model for the bulk crystallized polymer was discussed in the previous papers, where the dispersion behaviour of the specifically constructed model fairly well reproduced that of the crystallized polymer with various degrees of crystallinity. In this paper we will apply similar considerations to the oriented polymers.

Crystallized samples of polyethylene (PE), isotactic polypropylene (PP), polytetrahydrofuran (PTHF), polyethylene oxide (PEO) and polyoxymethylene (POM) were well annealed after an orientation was given by drawing or rolling. The higher order structure of these samples were surveyed by using x-ray small angle scattering. From the intensity distribution along equatorial, layer line or meridian, it was concluded that the microfibrils of these samples are united by heat-treatment to form huge fibrils with large diameter, and that the arrangement of the crystalline region (C) and the amorphous region (A) becomes regular in the way that A and C align in series along the fibre axis and in parallel along the direction perpendicular to the fibre axis. Figure 26 (a) shows the Hosemann’s model for the hot-drawn polyethylene. This model can be reasonably applied to the higher order structure of the other samples used here in view of the results of small angle x-ray scattering.

There is a general relationship common to all dispersion curves inspected here, i.e. those of the annealed oriented PE, PP, POM, PEO and PTHF,
Figure 18. Electron photomicrograph of single crystal of trans-1,4-polybutadiene prepared from benzene solution at $18^\circ$C and its electron diffraction pattern.
Figure 27. Temperature dependence of $E'$ and $E''$ along and perpendicular to the draw direction for the annealed samples of drawn polypropylene at 110 c/s. X-ray fibre diagram is for the same sample.

Figure 28. Temperature dependence of $E'$ and $E''$ in directions of 0, 20, 45, 70 and 90° to the draw direction for the annealed samples of drawn polytetrahydrofuran [The upper x-ray diffraction diagram is obtained with x-ray beam parallel to the drawing direction and the lower diagram is obtained with x-ray beam perpendicular to the draw direction (fibre diagram)]]
Figure 33. Rotational x-ray diffraction diagram of polytetraoxane crystal polymerized in a solid state by gamma-ray irradiation used for the viscoelastic measurements
that the modulus (tensile storage modulus at 110 c/s) along the fibre axis is higher than that of the perpendicular direction below the primary absorption temperature, they intersect at this temperature and above that temperature the modulus along the fibre axis becomes lower than the perpendicular one. Figures 27 and 28 (see art section) show the dispersion curves along and perpendicular to the draw direction for drawn annealed PP and PTHF respectively. X-ray diffraction patterns of the same samples are cited together. These are two examples of the general phenomena of modulus intersection. The primary absorptions are not distinct of linear polyethylene and polyoxymethylene and their intersecting temperatures are not clear, but they are considered to be located at around the temperatures of the latent primary absorptions \(^\text{16, 40}\).

The general relationship observed here will be qualitatively understood by adopting the Hosemann’s model Figure 26 (a) for the well annealed oriented polyethylene and by taking into account both the anisotropy of the polymer crystal and the approximately isotropic amorphous region. Figure 29 shows the schematic representation of the mechanical model corresponding

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\text{Figure 29. Schematic representations of dispersion curves of } E' \text{ (solid lines) in the parallel direction (a) and the perpendicular direction (b) to the draw direction. Calculations are made by use of the } E' \text{ values of the amorphous (A) and the crystalline region (C) as expressed by the broken lines. Part (c) represents the difference between the dispersion curves of the parallel direction and the perpendicular direction. Directions of applied force are indicated by the arrows in the model, which is a simplified one of Figure 26 (a).}
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MOTOWO TAKAYANAGI

with the Hosemann’s model of Figure 26 (a). The modulus of the crystalline region was assumed to be $10^{12}$ dyne/cm$^2$ for the parallel direction and $10^{10}$ dyne/cm$^2$ for the perpendicular direction, and the modulus of the amorphous region was assumed to be $10^{10}$ dyne/cm$^2$ below the glass transition temperature and $10^8$ dyne/cm$^2$ above the glass transition temperature. Figure 29 also shows the dispersion curves calculated for the cases when the force is applied in parallel and perpendicular to the draw direction. The model presented here is successful in qualitative explanation of the intersection of modulus, but as compared with the experimental results it is more reasonable to modify the AC series model like the one shown in Figure 26 (b). The modified model of Figure 26 (b) is improved in interpreting that the crystalline absorption can be clearly found along the draw direction and the modulus along the draw direction above $T_g$ does not decrease as much as expected from the simple AC series model. The modified model is composed of the laterally developed and oriented lamellar crystals, thin oriented crystals connecting these lamellae along the draw direction, and the inter-lamellar amorphous regions.

X-ray small angle scattering of the cold-drawn samples reveals the broad distribution of the microfibril diameter and the spatially irregular arrangement of the amorphous and the crystalline region. The modulus perpendicular to the fibre axis is expected to be largely decreased by the presence of microvoids between the microfibrils. The left-hand side of Figure 3

![Figure 30](image)

**Figure 30.** (a) Structural model of cold-drawn linear polyethylene presented by Hosemann, which is composed of splitted microfibrils with different diameters. (b) Mechanical model corresponding with the structural model in (a), expressing the various ways of response depending on the direction of applied force.

the model reproduced from Hosemann’s paper, which is given for the cold-drawn polyethylene. The right-hand side of Figure 30 is the corresponding mechanical model. Actually in the dispersion curves of cold-drawn linear polyethylene and polypropylene there could not be found any intersection of the moduli along the draw direction with that along the perpendicular direction as shown in Figures 31 and 32. Detailed inspection of the value of modulus leads us necessarily to adopt the anisotropy of the
amorphous region, which is caused by the insufficient relaxation of the oriented amorphous region due to low drawing temperature.

In order to understand the higher order structure of the industrial fibres or oriented films it would be very instructive to know the dispersion behaviour of the ideally oriented polymer crystal with the highest perfection. Single crystal of tetraoxane can be polymerized in a solid state by gamma-ray irradiation to form the polymer crystal composed of extended chains of polyoxymethylene. Figure 33 shows the rotational x-ray diffraction pattern of polytetraoxane crystal, which proves that molecular chains of polyoxymethylene are ideally oriented along the long axis of the crystal.
After extraction of residual monomer the tensile viscoelastic measurements were made along the long axis of the crystal. Figure 34 shows the tan δ vs. temperature curve for polytetraoxane (curve 1), polyoxymethylene fibre prepared by drawing the crystallized polymer (curve 2) and single crystal mats of polyoxymethylene (curve 3). Tan δ peak at about $-70^\circ$C corresponds to the secondary absorption and its mechanism of relaxation is ascribed to the local twisting motion of main chains in the defect region of crystal and in the frozen amorphous region. The fact that there is no secondary absorption in polytetraoxane crystal as shown in Figure 34 predicts that the polytetraoxane crystal is almost perfect, being absent from defect region. The single crystal sample (curve 3) in Figure 34 includes some such defect regions and the so-called fibre prepared by drawing (curve 2) has an unexpectedly large number of defects in its structure. Figure 35 shows the tensile storage modulus and the tensile loss modulus of the three samples as a function of temperature. The storage modulus of polytetraoxane displays an outstanding constancy of the modulus value very close to the melting temperature $190^\circ$C, while the modulus value of drawn polyoxymethylene (broken lines) decreases gradually from about the secondary absorption temperature, suggesting many kinds of relaxation mechanisms associated with the various types of disorder. Referring to the dispersion curves of the three samples, the drawn fibre rather resembles in some point the single crystal sample. This suggests that the lamellar structure composed of folded molecules has a possibility of being included in the oriented fibre. To prove it, however, it is necessary to have more information on fibre structure.
CRYSTALLIZED STATE OF POLYMER IN ITS DISPERSION BEHAVIOUR

Figure 35. $E'$ and $E''$ vs. temperature for three kinds of samples of polyoxymethylene [●] is for the polytetraoxane crystal, (— — —) is for the drawn sample, and (○) is for the single crystal mat. These samples are the same as in Figure 34.

Measurement of viscoelastic absorption is one of the methods of obtaining the information on molecular motions. The dielectric absorption and the broad line n.m.r. absorption are also available. The purpose of this article is to draw attention to the fact that the information from dispersion behaviour is useful to approach to the true features of the crystallized polymer by combined use of the morphological and of the structural methods.

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MOTOWO TAKAYANAGI

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