Uniaxial and Mixed Orientations of Poly(ethylene oxide) in Nanoporous Alumina Studied by X-ray Pole Figure Analysis

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ABSTRACT:

The orientation of polymers under confinement is a basic, yet not fully understood phenomenon. In this work, the texture of poly(ethylene oxide) (PEO) infiltrated in nanoporous anodic alumina oxide (AAO) templates was investigated by X-ray pole figures. The influence of geometry and crystallization conditions, such as pore diameter, aspect ratio, and cooling rates, was systematically examined. All the samples exhibited a single, volume-dependent crystallization temperature ($T_c$) at temperatures much lower than that exhibited by bulk PEO, indicating “clean” microdomains without detectable heterogeneous nucleation. An “orientation diagram” was established to account for the experimental observations. Under very high cooling rates (quenching), crystallization of PEO within AAO was nucleation-controlled, adopting a random distribution of crystallites. Under low cooling rates, growth kinetics played a decisive role on the crystal orientation. A relatively faster cooling rate (10 °C/min) and/or smaller pores lead to the $<120>\parallel$ pore axis ($\vec{n}$) mode (uniaxial orientation). When the cooling rate was lower (1 °C/min), and/or the pores were larger, a mixed orientation, with a coexistence of $<120>\parallel\vec{n}$ and $<010>\parallel\vec{n}$, was observed. The results favor the kinetic model where the fastest growth direction tends to align parallel to the pore axis.

1. INTRODUCTION

The crystallization behavior of polymers under confinement are attracting increasingly attention because it provides a model system to understand basic questions about polymer nucleation and growth kinetics, as well as its close relevance to modern nanotechnology, such as nanofabrication and functional nanodevices.1,2,3,4,5,6,7 Nanoporous anodic alumina oxide (AAO) templates have been widely used to construct a uniform confined environment since the development of electrochemical method to fabricate well-ordered arrays.8 The nucleation and crystallization of polymers within AAO templates has been the subject of several reviews.2,9,10

Two general observations have been reported for infiltrated polymers within AAO, i.e., altering of the
nucleation mechanism and anisotropic crystal growth. Much larger supercoolings are observed for polymers infiltrated in AAO, indicating a change of nucleation mechanism from heterogeneous to homogenous or surface nucleation.\textsuperscript{11,12} The presence of fractionated crystallization in infiltrated polymers in AAO templates, where several crystallization peaks are observed upon cooling from the melt\textsuperscript{5} has been recently assigned to the percolation of surface film residues that have not been completely removed during the cleaning procedure performed after infiltration.\textsuperscript{13}

Crystal orientation within AAO nanopores is another interesting phenomenon that has also attracted much attention. Polymer chains within the crystals typically align with their chain axis (i.e., $c$ axis) perpendicular to the pore axis (defined as $c \perp \vec{n}$, hereafter).\textsuperscript{14} This has been explained by a “kinetic effect” by Steinhart et al.,\textsuperscript{15} i.e., those crystal lamellae with growing direction not parallel to the pore axis are blocked by the wall, while the crystal lamellae with the $<hk0>^*$ direction parallel to the pore axis are free to grow. It has been realized that the orientation mode of “$c \perp \vec{n}$” is a very “weak” restriction for crystal growth, usually resulting in a low degree of anisotropy.\textsuperscript{13}

A geometrically similar 2D confined system is the well-studied cylinder-forming block copolymers, however, the orientation model was quite different with respect to the “$c \perp \vec{n}$” mode.\textsuperscript{16} Huang et al. studied the crystal orientation of PEO block nanocylinders (with a diameter of 13.7 nm) within a PS matrix in a PEO-$b$-PS/PS blend.\textsuperscript{17} It was shown that the crystallization temperature played a significant role in the crystal orientation. At very low crystallization temperature ($T_c$), PEO crystals were randomly oriented within the confined cylinders. Upon increasing the crystallization temperature ($-30 \textdegree C \leq T_c \leq 0 \textdegree C$), the crystal orientation changed to be inclined with respect to the cylinder axis. When the sample was crystallized at $T_c \geq 2\textdegree C$, a uniaxial orientation with the $<120>^*$ direction parallel to the pore axis was found. This orientation mode resulted from the alignment of maximum PEO crystal growth direction ($<120>^*$) with the channel axis.

Table 1 summarizes the orientation of polymer crystals confined in AAO nanopores. Except for several
particular reports,\textsuperscript{12,15,18,19} the majority of the systems exhibit uniaxial orientation or a mixed structure with two types of uniaxial orientations. It is noticed that the structure of many systems could be explained by the alignment of the fastest growth direction with the pore axis, such as in PEO\textsuperscript{20,21,22} and in PE.\textsuperscript{23,24} Two modes of crystal orientation were identified in PEO infiltrated within AAO, namely, perpendicular orientation with the normal direction of (120) plane (i.e., \textit{<120>\textsuperscript{*}} direction) aligning parallel to the pore axis and tilt orientation with the \textit{<120>\textsuperscript{*}} tilted 45° away from the pore axis.\textsuperscript{20,21} Two types of orientation were also observed in poly(caprolactone) (PCL),\textsuperscript{13} isotactic polypropylene (iPP),\textsuperscript{13} PVDF,\textsuperscript{25,26} and syndiotactic polypropylene (sPP).\textsuperscript{27} There is still a lack of a reasonable explanation of the coexistence of different orientation modes.

Thermodynamics or surface effects play a role in the crystal orientation as well. When the size of the AAO pores is smaller than the contour length of the low molecular weight poly(ethylene oxide) (PEO), the polymer crystallized into its thermodynamic stable structure by maintaining the extended chain crystal with polymer chains align parallel to the AAO axis.\textsuperscript{22} The “parallel orientation” observed in infiltrated syndiotactic polystyrene (sPS) within AAO templates was explained by a surface nucleation effect.\textsuperscript{18}

Apart from the lack of a general model for the orientation of polymers confined in AAO templates, it should be noted that the characterization of the texture is frequently inadequate in the literature. The most commonly applied two-dimensional X-ray diffraction method, only provides limited information of the reciprocal space. Further, the influence of the pore diameter, aspect ratio, as well as crystallization conditions have not been quantitatively investigated yet. In this work, X-ray pole figure measurements are employed to characterize infiltrated PEO in AAO templates to get a full picture of the crystal orientation. It was shown that PEO predominately adopted a uniaxial orientation in AAO with the \textit{<120>\textsuperscript{*}} direction parallel to the pore axis ($\vec{n}$). Smaller diameter and higher aspect ratio resulted in higher degrees of orientation. Cooling rate played the most significant role in the crystal orientation. A “tilted orientation” of PEO was confirmed, which is in line with the
However, a new explanation was proposed to justify the multiple uniaxial orientational phenomena by investigating the influence of geometry and crystallization conditions.

### Table 1. Summary of the Orientation of Infiltrated Polymer Crystallites within AAO Nanopores.

<table>
<thead>
<tr>
<th>Polymer</th>
<th>Infiltration method</th>
<th>Surface layer$^a$</th>
<th>Crystallization condition</th>
<th>Orientation$^a$</th>
<th>$\Phi_{AAO}$ (nm)</th>
<th>Ref.</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>PEO</strong></td>
<td>melt, nanorod</td>
<td>N</td>
<td></td>
<td></td>
<td>30, 60 &amp; 100</td>
<td></td>
</tr>
<tr>
<td></td>
<td>solution, nanotube</td>
<td>N</td>
<td>-40 °C, -20 °C, 0 °C, &amp; 20 °C</td>
<td>$&lt;120&gt;^*$ $\parallel$ $\bar{n}$ or tilted 45°</td>
<td>23 &amp; 89</td>
<td>20</td>
</tr>
<tr>
<td></td>
<td>melt, nanorod</td>
<td>N</td>
<td>-40 °C, -20 °C, 0 °C, &amp; 20 °C</td>
<td>$&lt;120&gt;^*$ $\parallel$ $\bar{n}$ or tilted 45°</td>
<td>23 &amp; 89</td>
<td>21</td>
</tr>
<tr>
<td></td>
<td>melt</td>
<td>N</td>
<td></td>
<td>$c \perp \bar{n}$</td>
<td>20</td>
<td>12</td>
</tr>
<tr>
<td></td>
<td>melt, nanotube</td>
<td>N</td>
<td>$&lt;120&gt;^*$ $\parallel$ $\bar{n}$</td>
<td>400</td>
<td>19</td>
<td></td>
</tr>
<tr>
<td><strong>PE</strong></td>
<td>melt, nanorod</td>
<td>N</td>
<td>1 °C/min</td>
<td>mainly $\bar{b} \parallel \bar{n}$</td>
<td>60</td>
<td>23</td>
</tr>
<tr>
<td></td>
<td>melt, nanorod</td>
<td>N</td>
<td>0.5 °C/min</td>
<td>$\bar{b} \parallel \bar{n}$</td>
<td>15 to 220</td>
<td>24</td>
</tr>
<tr>
<td></td>
<td>melt, nanorod</td>
<td>Y</td>
<td>120 °C</td>
<td>mainly $\bar{b} \parallel \bar{n}$</td>
<td>200</td>
<td>28</td>
</tr>
<tr>
<td></td>
<td>melt, nanotube</td>
<td>N</td>
<td>$\bar{b} \parallel \bar{n}$</td>
<td>400 &amp; 1000</td>
<td>14</td>
<td></td>
</tr>
<tr>
<td><strong>PVDF</strong></td>
<td>melt, nanotube</td>
<td>N</td>
<td>1 °C/min, 136 °C, 140 °C or 152 °C</td>
<td>$&lt;020&gt;^<em>$ or $&lt;110&gt;^</em> \parallel \bar{n}$</td>
<td>400</td>
<td>25</td>
</tr>
<tr>
<td></td>
<td>melt, nanotube</td>
<td>N</td>
<td>1 °C/min</td>
<td>$&lt;020&gt;^<em>$ or $&lt;110&gt;^</em> \parallel \bar{n}$</td>
<td>400</td>
<td>26</td>
</tr>
<tr>
<td></td>
<td>solution, nanorod</td>
<td>Y</td>
<td>60 °C</td>
<td>$\bar{b} \parallel \bar{n}$</td>
<td>150-200</td>
<td>29</td>
</tr>
<tr>
<td><strong>P(VDF-TrFE)</strong></td>
<td>melt, nanorod &amp; nanotube</td>
<td>N</td>
<td>2 °C/min</td>
<td>$&lt;200&gt;^<em>$ or $&lt;110&gt;^</em> \parallel \bar{n}$</td>
<td>400</td>
<td>30</td>
</tr>
<tr>
<td></td>
<td>solution, nanorod</td>
<td>Y</td>
<td>135 °C</td>
<td>$a \parallel \bar{n}$</td>
<td>40, 60 &amp; 80</td>
<td>31</td>
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<tr>
<td></td>
<td>melt, nanorod</td>
<td>Y</td>
<td>125 °C</td>
<td>$c \perp \bar{n}$</td>
<td>15 &amp; 40</td>
<td>32</td>
</tr>
<tr>
<td></td>
<td>melt, nanotube</td>
<td>Y</td>
<td>125 °C</td>
<td>random</td>
<td>200</td>
<td></td>
</tr>
<tr>
<td><strong>sPP</strong></td>
<td>melt, nanorod</td>
<td>N</td>
<td>slow cooling</td>
<td>$&lt;020&gt;^*$ $\parallel \bar{n}$</td>
<td>110, 300</td>
<td>27</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td>$&lt;200&gt;^<em>$ or $&lt;020&gt;^</em> \parallel \bar{n}$</td>
<td>30</td>
<td></td>
</tr>
<tr>
<td><strong>iPP</strong></td>
<td>melt, nanorod</td>
<td>N</td>
<td>random (200 nm)</td>
<td>$a^* \parallel \bar{n}$ (mixed)</td>
<td>15, 40</td>
<td>33</td>
</tr>
<tr>
<td></td>
<td>melt, nanorod</td>
<td>N</td>
<td>1 °C/min</td>
<td>$a^* \parallel \bar{n}$ (mixed)</td>
<td>20 to 400</td>
<td>13</td>
</tr>
<tr>
<td><strong>PCL</strong></td>
<td>melt, nanorod</td>
<td>N</td>
<td>1 °C/min</td>
<td>$a^* \parallel \bar{n}$ (mixed)</td>
<td>20 to 400</td>
<td>13</td>
</tr>
<tr>
<td></td>
<td>melt, nanorod</td>
<td>N</td>
<td>1 °C/min</td>
<td>$\Phi_{AAO} \leq 40$ nm, $c \perp \bar{n}$</td>
<td>25, 35, 65 &amp; 200</td>
<td>34</td>
</tr>
<tr>
<td></td>
<td>melt, nanorod</td>
<td>N</td>
<td>3 °C/min, or 50 °C/min / 0 °C/min</td>
<td>$&lt;110&gt;^<em>$ or $&lt;100&gt;^</em> \parallel \bar{n}$</td>
<td>200</td>
<td></td>
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<tr>
<td><strong>sPS</strong></td>
<td>melt, nanorod</td>
<td>Y</td>
<td>255 °C</td>
<td>$c \perp \bar{n}$</td>
<td>32, 80 &amp; 200</td>
<td>18</td>
</tr>
<tr>
<td></td>
<td>melt, nanorod</td>
<td>N</td>
<td>260 °C</td>
<td>$c \perp \bar{n}$ (b-form)</td>
<td>80 &amp; 200</td>
<td>35</td>
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<tr>
<td><strong>iPS</strong></td>
<td>melt, nanorod</td>
<td>Y</td>
<td>170 °C</td>
<td>$&lt;100&gt;^* \parallel \bar{n}$</td>
<td>300</td>
<td>36</td>
</tr>
<tr>
<td></td>
<td>melt, nanorod</td>
<td>Y</td>
<td>$&lt;110&gt;^<em>$ or $&lt;100&gt;^</em> \parallel \bar{n}$</td>
<td>65</td>
<td></td>
<td></td>
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<tr>
<td><strong>PLLA</strong></td>
<td>melt, nanorod</td>
<td>N</td>
<td>$&lt;110&gt;^<em>$ or $&lt;200&gt;^</em> \parallel \bar{n}$</td>
<td>40, 75, 200, &amp; 385</td>
<td>38</td>
<td></td>
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<tr>
<td><strong>PBA</strong></td>
<td>melt, nanotube</td>
<td>N</td>
<td>20 °C/min</td>
<td>mainly $\bar{b} \parallel \bar{n}$ (b-phase)</td>
<td>30 &amp; 70</td>
<td>39</td>
</tr>
<tr>
<td></td>
<td>melt, nanotube</td>
<td>N</td>
<td>33 °C or 35 °C</td>
<td>mainly $\bar{b} \parallel \bar{n}$ (α-phase)</td>
<td>100 &amp; 200</td>
<td>40</td>
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<tr>
<td><strong>P3HT</strong></td>
<td>solution, nanotube</td>
<td>N</td>
<td>as-prepared</td>
<td>$\bar{b} \parallel \bar{n}$ (majority) or $a \parallel \bar{n}$</td>
<td>240</td>
<td>41</td>
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<tr>
<td>200 °C</td>
<td>10 °C/min</td>
<td>b∥(\mathbf{n}) (majority) or a∥(\mathbf{n})</td>
<td></td>
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<td></td>
<td></td>
</tr>
<tr>
<td>as-prepared</td>
<td>b∥(\mathbf{n})</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>annealed at 200 °C</td>
<td>b∥(\mathbf{n})</td>
<td></td>
<td></td>
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</tr>
<tr>
<td>10 °C/min</td>
<td>b∥(\mathbf{n}) (majority) or a∥(\mathbf{n})</td>
<td></td>
<td></td>
<td></td>
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<td></td>
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</table>

<table>
<thead>
<tr>
<th>Solution, Nanotube</th>
<th>PTT</th>
<th>Melt, Nanorod</th>
<th>Y</th>
<th>3 °C/min</th>
<th>a∥(\mathbf{n})</th>
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<tbody>
<tr>
<td>N</td>
<td>Y</td>
<td>0.5 °C/min</td>
<td>c∥(\mathbf{n})</td>
<td></td>
<td></td>
</tr>
<tr>
<td>N</td>
<td>Y</td>
<td>50 °C/min</td>
<td>&lt;530&gt;<em>∥(\mathbf{n}) (majority) or &lt;200&gt;</em>∥(\mathbf{n})</td>
<td></td>
<td></td>
</tr>
<tr>
<td>N</td>
<td>low cooling rates</td>
<td>&lt;008&gt;*∥(\mathbf{n})</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>nylon-12</td>
<td>melt, Nanorod</td>
<td>N</td>
<td>160 °C</td>
<td>b⊥(\mathbf{n})*</td>
<td></td>
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</tbody>
</table>

*a* The letter “N” is an abbreviation for “No” and “Y” is an abbreviation for “Yes”.

*b* \(<hkl>*\) indicates the direction in reciprocal space, which is normal to the \((hkl)\) plane in real space. The \(\mathbf{n}\) represents the pore axis, which is normal to the surface plane of the AAO template.

*c* The chain axis is defined as \(b\)-axis in nylon-12.

2. EXPERIMENTAL SECTION

2.1 Materials and Sample Preparation

Close to monodisperse poly(ethylene oxide) (PEO), with number-average molecular weight of 10,000 g/mol and polydispersity of 1.05, was purchased from Polymer Source Inc. The AAO templates with pore diameters of 40, 100 and 400 nm, and pore lengths of 100, 50, 20 and 5 μm were prepared by a two-step electrochemical anodization of aluminum as described in our previous paper.\(^\text{13}\)

Before infiltration with PEO solution, the AAO templates were washed with acetone and ethanol to remove possible impurities. The concentration of the PEO chloroform solution was 10 mg/ml. The solution was drop casted on the AAO templates and dried in air. Then the AAO templates were treated at 100 °C under vacuum for 1 hour to remove residual solvent. The casting-heating process was repeated several times to ensure that the nanopores were fully infiltrated. The residual PEO on the surface of AAO templates was removed using a polish cloth to ensure the separate crystallization of PEO within the nanopores. Unless specified, the samples were crystallized at a constant cooling rate of 10 °C/min.

2.2 Characterization

Thermal analysis was conducted with a differential scanning calorimeter (DSC Q2000, TA). The instrument
was calibrated with indium before measurements. The samples of ~5 mg with the aluminum base were weighed, and encapsulated in aluminum pans for DSC measurements. All the samples were first heated to 100 °C and held for 3 min to eliminate the thermal history. Then the samples were scanned from 100 to -50 °C at a cooling and heating rate of 10 °C/min under high purity nitrogen atmosphere.

2D Wide angle X-ray diffraction (WAXD) measurements were carried out at room temperature on a Xeuss 2.0 SAXS/WAXS system (Xenocs SA, France). CuKα X-ray source (GeniX3D Cu ULD), generated at 50 kV and 0.6 mA, was utilized to produce X-ray radiation with a wavelength of 1.5418 Å. A semiconductor detector (Pilatus 300 K, DECTRIS, Swiss) with a resolution of 487 × 619 pixels (pixel size = 172 × 172 μm²) was used to collect the scattering signals. The scattering geometry is indicated in Figure 1. The ψ angle and φ angle are all set to 0°. The X-ray irradiates the sample along the x-axis with an incident angle of 3° with respect to the AAO surface. The 2D detector is placed perpendicular to the incident beam. The exposure time for each pattern is 5 min. The one-dimensional intensity profiles were integrated from the 2D WAXS patterns, averaged along the azimuthal angle.

The pole figures were measured with D8 DISCOVER X-ray diffractometer (Bruker) in reflection geometry. The wavelength of the radiation was 1.5406 Å. A 1D detector, the Bruker LYNXEYE, was employed to collect the intensity. Prior to the pole figure measurement, a specular scan was carried out to confirm the 2θ angle of the reflections. The diffraction angles were set as 19.1° and 23.3° for the two strongest reflection peaks (120) and (112)/[(032)/(132)/(212)], respectively. These crystal planes have similar d-spacing of about 0.39 nm. Figure 1 shows a schematic of the measurement geometry. The sample was mounted on top of a sample stage on an Eulerian cradle. During measurement, the sample rotated stepwise around the z (n) axis (φ angle) and the x axis (ψ angle). The range of ψ was set from 0° to 80° with an angular interval of 5° and the exposure time of 10 s.
3. RESULTS AND DISCUSSION

3.1 DSC and WAXD

Figure 2. (a) DSC cooling curves of infiltrated PEO in AAO templates with different pore size. (b) Crystallization temperature ($T_c$, black square), calculated $T_c$ (blue sphere) and melting temperature ($T_m$, red triangle) of infiltrated PEO in AAO as a function of pore volume. (c) 2D WAXD pattern of PEO confined within the AAO template with pore depth of 100 μm, and pore diameter of 100 nm. The beam center is represented by a white cross. (d) Corresponding 1D intensity profile of the infiltrated PEO, obtained by averaging the intensity along the azimuthal angle.
Figure 2a shows that all the samples crystallized at very large supercoolings, which agrees with previous reports. Only one peak is observed for all the samples, indicating that the number of nanopores without any active nucleating heterogeneity for PEO is much larger than the number of such heterogeneities in bulk PEO. Therefore, the overwhelming majority of the nanopores did not contain heterogeneous nucleation sites and no surface film that could percolate the pores was formed in any of the infiltrated AAO templates. The crystallization/melting temperatures, as a function of pore volume, are plotted in Figure 2b. A very clear trend can be observed, that is, the crystallization temperature of the samples decreases with pore volume. The estimated $T_c$ according to the empirical formula obtained by Müller et al. for homogenously nucleated PEO phases within block copolymers and for PEO droplets, $T_c(°C) = -41.8 + 2.89 \log(V_d (\text{nm}^3))$, was also plotted in Figure 2b. The $T_c$ values differ only about 2 ~ 4 °C. Therefore, as previously reported for PEO, homogeneous nucleation probably occurs inside the volume of infiltrated material without any influence of the alumina surface. The $T_m$ values are nearly constant, indicating the crystals have similar thermodynamic stability, which agrees with previous reports.

A typical 2D WAXD pattern of infiltrated PEO is shown in Figure 2c and the corresponding intensity profile is plotted in Figure 2d. The (120) reflection locates on the meridian and the (112)/(032)/(132)/(212) reflection locates mainly on the off-meridian area. The basic features of the 2D pattern agree well with the previous reports. Although the pattern only covers a slice of the reciprocal space depending on the scattering vector, such pattern has been interpreted as an evidence for the $<120>^*$ direction of the PEO crystals aligning parallel to the pore axis ($<120>^* \parallel \vec{n}$).

### 3.2 Pole Figures of Infiltrated PEO

The pole figures corresponding to (120) and (112)/(032)/(132)/(212) reflections of the infiltrated PEO crystals in AAO templates with different diameters are shown in Figure 3. The depth of the templates (100 μm) and the
thermal history of the samples (10°C/min cooling) are the same. The intensity of the (120) pole figures show maximum values at \( \psi = 0^\circ \), which means that the (120) crystal plane normal is preferentially aligned parallel to the pore axis. The pole figures of (112)/(032)/(1\overline{3}2)/(\overline{2}12) reflections resemble rings with maxima located at \( \psi \approx 30^\circ \).

As shown in Table 2, the crystal planes with similar \( d \)-spacings, occasionally, have similar inclination angle with respect to the (120) plane within the range of 34-35°. Therefore, the pole figures agree with the texture that the (120) plane normal or the \(<120>*\) in reciprocal space align with the pore axis. The (120) crystal plane is the fastest growth plane in PEO.\(^{49,50,51,52}\) Therefore, the commonly reported result that the maximum growth direction aligns along the pore axis is confirmed, which agrees with the texture of PEO crystals in strongly segregated block copolymers crystallized at low supercoolings.\(^{17}\)

Table 2: The Angle between Different Crystal Planes in PEO.

<table>
<thead>
<tr>
<th>plane</th>
<th>120</th>
<th>112</th>
<th>032</th>
<th>1\overline{3}2</th>
<th>\overline{2}12</th>
<th>100</th>
<th>010</th>
</tr>
</thead>
<tbody>
<tr>
<td>( d )-spacing (Å)</td>
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<td>23.0</td>
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<td>35.1</td>
<td>33.9</td>
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<tr>
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<td>28.7</td>
<td>29.4</td>
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The influence of pore diameter is examined first, for which the templates have the same depth and the same thermal history. The general feature of those pole figures are similar. The orientation condition can be seen more clearly in the intensity profiles along \( \psi \) obtained by averaging circularly along \( \varphi \) angles (see Figure 4). It is shown that, as pore diameter decreases, the degree of orientation remains very similar. This is in accordance with our previous study,\(^{22}\) where the orientation of the crystal planes basically remains the same as the pore diameter is varied above a critical value. Another noteworthy feature is that a shoulder peak at \( \psi = 45^\circ \) appeared for the (120) reflection for the sample with 400 nm AAO (arrow in the figure).

The tilted (120) orientation has been observed recently by Liu and Chen,\(^{20,21}\) where they explained it by the possible “double constrains” imposed by the AAO wall and by the neighboring crystallites when the nucleation
density is high. Although apparently possible, however, this model could not fully explain why the tilt angle (45°) was independent of crystallization temperatures (i.e., nucleation densities), molecular weights, and pore diameters, all of which are expected to influence the “strength” of the constricts caused by the neighboring crystallites. The tilt angle seems more like an intrinsic property of the PEO crystal rather than an environmental or kinetic effect. To account for the above observations, therefore, we propose an alternative explanation. Table 2 lists the angles between different crystal planes in PEO, calculated according to the unit cell parameters proposed by Takahashi and coworkers.\textsuperscript{53} It is interesting to find that the angle between (120) plane and (100) or (010) plane happened to be 45°. According to crystal growth studies on PEO, it has been reported that the (010) crystal plane is another fast growth plane of PEO.\textsuperscript{50,51,52} If a minor PEO crystal population assembles within the AAO templates with the \langle010\rangle* parallel to the pore axis (\langle010\rangle* \parallel \vec{n} ), the reflection of (120) plane of those crystals will exhibit a 45° tilt angle with respect to the pore axis, thereby satisfactorily explaining the observed results.

\begin{figure}[h]
\centering
\includegraphics[width=\textwidth]{pole_figures.png}
\caption{Pole figures of (a - c) the (120) reflections, and (d - f) the (032) reflections of infiltrated PEO in AAO templates with pore depth of 100 μm, and pore diameters of (a, d) 400 nm, (b, e) 100 nm, and (c, f) 40 nm.}
\end{figure}
Azimuthal profiles of (a) the (120) reflections, and (b) the (032) reflections of infiltrated PEO in AAO templates with pore depth of 100 μm, and pore diameters of 400 nm, 100 nm, and 40 nm.

3.3 Effect of Pore Depth

There are few studies concerning the effect of pore depth which influences the volume and shape of the pore and may influence the crystal orientation as well. In extreme cases when the pore depth is very small, a transition from 2D to 3D confinement would occur and the crystallites would be statistically random. Figure 5 shows the pole figures of PEO crystal planes confined in AAO with the same diameter (100 nm) and different pore depths (100, 50, 20, and 5 μm), while the azimuthal profiles are shown in Figure 6.

Before the pole figure measurements, the samples were first held at 100 °C for 3 min to erase their thermal history, and then crystallized at a cooling rate of 10 °C/min. The features of the pole figures are very similar, exhibiting predominately the \(<120>*\parallel \vec{n}\) texture. The azimuthal profiles indicate that the degree of orientation increases with pore depth and thus confirms our hypothesis, although the smallest depth of the pores (5 μm) is still
sufficiently large as compared with the diameter (100 nm).

**Figure 5.** Pole figures of (a - d) the (120) reflections, and (e - h) the (032) reflections of infiltrated PEO in AAO templates with pore diameter of 100 nm, and pore depths of (a, e) 100 μm, (b, f) 50 μm, (c, g) 20 μm, and (d, h) 5 μm.

**Figure 6.** Azimuthal profiles of the (120) reflection (a) and the (032) reflection (b) of infiltrated PEO in AAO templates with pore diameter of 100 nm, and pore depths of 100 μm, 50 μm, 20 μm, and 5 μm.
3.4 Effect of Cooling Rate

It could be clearly seen in Figure S1 (supporting information) that the crystallization temperature decreased with the increase of cooling rate, and only one crystallization peak was observed for all the cooling rates. Figure 7 shows the pole figures of infiltrated PEO in AAO templates with different cooling rates. Once more, the general feature of the orientation mode is the same. However, the degree of orientation significantly decreases when the sample is quenched in liquid nitrogen. This is more pronounced in the azimuthal profiles of the pole figures (see Figure 8).

The cooling rate influences the interplay between nucleation rate and crystal growth rate, therefore, it has a strong influence on the developed crystalline texture. Since the PEO nucleates homogeneously within AAO, at lower cooling rates or higher isothermal crystallization temperatures, nucleation is the dominating step and crystal growth proceeds almost instantaneously. Under this circumstance, the crystallization kinetics would be first order with an Avrami index of 1.9.54 With the increase of cooling rate or decrease of crystallization temperature, the nucleation rate will increase and the growth rate will decrease relative to one another. Under the extreme condition of ultra-high nucleation rate, many nuclei will form simultaneously within in one pore and the crystals grow little before they impinge with adjacent crystals. In this case, the nuclei will not “feel” the restricted geometry, hence isotropic texture would be observed. This is most obvious in the quenched sample.

![Pole figures](image)

**Figure 7.** Pole figures of (a - c) the (120) reflections, and (d - f) the (032) reflections of infiltrated PEO in AAO
template with the pore diameter of 100 nm, and pore depth of 100 μm, crystallized at different cooling rate. (a, d) crystallized at a cooling rate of 1 °C/min, (b, e) crystallized at a cooling rate of 10 °C/min, and (c, f) quickly quenched into liquid nitrogen.

Figure 8. Azimuthal profiles of (a) the (120) reflections, and (b) the (032) reflections of infiltrated PEO in AAO template with the pore diameter of 100 nm, and pore depth of 100 μm, crystallized at different cooling rate.

A closer look at Figure 8a shows that a small peak appears at ψ = 45° in the azimuthal profile of the (120) reflection for the PEO infiltrated in the AAO template with a cooling rate of 1 °C/min. The fact that a lower cooling rate is beneficial for the <010>*∥n⃗ orientation probably indicates the different temperature dependence of the growth rates of different planes. It is probable that at a lower cooling rate, the growth rate of (010) plane is comparable to that of (120) plane. Thus a mixed structure of <120>*∥̅n⃗ and <010>*∥̅n⃗ can be observed. Marentette et al. have shown in bulk PEO spherulites that the dominant crystal growth face transformed from (120) plane at lower supercoolings (Tc > 51°C) to (010) plane at higher supercoolings (Tc < 51°C). However, no crystal
growth data is available at extremely high supercoolings ($T_c < -10^\circ$C).

Figure 9 shows the azimuthal profiles of the (120) pole figures of other samples. The PEO infiltrated in AAO with a diameter of 100 nm and a depth of 50 μm (Figure 9a) exhibits similar features as in Figure 8. However, the $\psi = 45^\circ$ peak is not observed at all, even at the lowest cooling rate (1°C/min), for the sample with a smaller depth (20 nm, Figure 9b). For the sample with 400 nm diameter and 100 μm length, the $\psi = 45^\circ$ is observed for both 1 and 10°C/min. The infiltrated PEO within AAO with 40 nm diameter and 100 μm length does not show the $\psi = 45^\circ$ (Figure 9d).

![Figure 9. Azimuthal profiles of the (120) reflections of infiltrated PEO in AAO template with the pore diameter of 100 nm, and pore depths of (a) 50 μm and (b) 20 μm, and in AAO template with the pore depth of 100 μm, and pore diameters of (c) 400 nm, and (d) 40 nm, crystallized at different cooling rate.](image-url)
The pore diameters and pore depths of the AAO templates were 400 nm - 100 μm (square), 100 nm - 100 μm (circle), 100 nm - 50 μm (down triangle), 100 nm - 20 μm (diamond), 40 nm - 100 μm (up triangle), respectively.

The above results can be illustrated in an “orientation diagram” as shown in Figure 10. The diagram can be divided into three regions. Under super-high cooling rates (such as quenching), the nucleation density is extremely high, resulting in a random orientational texture, irrelevant to the size of the domains. When the cooling rates are lower, the texture of the samples are determined by kinetic effects, i.e., the fastest growth direction aligns parallel with the pore axis. Two textures are found at low cooling rates. In the top-left region with smaller pores and/or higher cooling rates, only the $<120>^* || \vec{n}$ mode is observed. The bottom-right region with larger pores and/or lower cooling rates exhibits mixed orientation modes of $<120>^* || \vec{n}$ and $<010>^* || \vec{n}$.

**3.5 Remarks on the Orientation Model**

Our results show that the orientation of infiltrated PEO in AAO templates agrees largely with the report of Huang et al.\textsuperscript{17} in cylinder-forming block copolymers containing PEO, which is different from the model proposed by Steinhart et al.\textsuperscript{15} in PVDF. Which model is more probable? A first look at Table 1 tells us that most of the reports fit the uniaxial/mixed orientation model. Probably, the PVDF is the only polymer where all $(hk0)$ planes are able to grow under confinement. On the other hand, surface film residues of infiltrated polymers often manifest itself by showing multiple crystallization peaks.\textsuperscript{13} It is not clear how significant the residual polymer films on the surface of
the AAO templates would influence the orientation of crystals.

Regardless of incompleteness, it is fairly safe to state “the chain axis is perpendicular to the pore axis”. This allows the crystals to “rotate freely” around the chain axis, leading to the texture with all the (hk0) visible on the specular scan of XRD. Meanwhile, the crystals could also rotate freely around the pore axis. Except under super-high cooling rates, this statement seems always valid for crystallization under confinement governed by kinetics.

Another implication of the present work is that the crystal growth kinetics under very high supercoolings can be probed by studying the orientation feature of polymer confined in AAO, which is not possible for bulk samples. At least, the fastest growth plane could be readily read out. An important assumption is that the primary nuclei are randomly oriented. This is most likely true for homogeneous nucleation, still need to be explored in systems with surface nucleation.

4. CONCLUSIONS

In this work, a semi-quantitative study of the orientation of PEO confined in AAO templates was carried out for the first time by pole figure measurements. The influence of pore diameter, pore length and cooling rate was examined. Uniaxial orientation and mixed orientations were observed in PEO, in favor of the kinetic model proposing that the fastest growth direction aligns parallel to the pore axis. An “orientation diagram” was established to account for the two most important factors governing the crystal orientation texture. By quenching, crystallization of PEO within AAO was nucleation-controlled, exhibiting a random distribution of crystallites. Under low cooling rates, crystal growth rate played a decisive role on the crystal orientation. Relatively faster cooling rates and smaller pores lead to the $\langle 120\rangle^* \parallel \vec{n}$ orientation. When the cooling rate was even lower, and/or the pores were larger, mixed orientations, with a coexistence of $\langle 120\rangle^* \parallel \vec{n}$ and $\langle 010\rangle^* \parallel \vec{n}$, was observed. The texture of the polymer within AAO provides hints on the crystal growth kinetics at high supercoolings which are
inaccessible in bulk studies.

ACKNOWLEDGEMENTS

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Supporting Information

Hermans’ orientation parameter; DSC cooling curves of the infiltrated PEO in AAO template with different cooling rates; Crystallization temperature as a function of cooling rate.

REFERENCES


24 Shin, K.; Woo, E.; Jeong, Y. G.; Kim, C.; Huh, J.; Kim, K.-W. Crystalline Structures, Melting, and Crystallization of


34 Suzuki, Y.; Duran, H.; Akram, W.; Steinhart, M.; Floudas, G.; Butt, H. J. Multiple Nucleation Events and Local


44 Martin, J.; Scaccabarozzi, A. D.; Nogales, A.; Li, R. P.; Smilgies, D. M.; Stingelin, N. Confinement Effects on the


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Uniaxial and Mixed Orientations of Poly(ethylene oxide) in Nanoporous Alumina Studied by X-ray Pole Figure Analysis

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