

ORIGINAL ARTICLE

Development of a fiber structure in poly(vinylidene fluoride) by a CO₂ laser-heated drawing process

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Rapid and uniform heating by CO₂ laser radiation can fix the position where necking occurs. Therefore, this study investigated the development of a fiber structure in poly(vinylidene fluoride) in continuous drawing by *in situ* measurement using synchrotron X-ray radiation with a time resolution of several hundred microseconds. Two neck-deformation behaviors were observed in the laser drawing: a moderate neck deformation under low drawing stress and a steep neck deformation under high drawing stress. The low drawing stress resulted in a mixture of α - and β -crystals in which the β -crystal was formed within 1 ms after the necking, earlier than the α -crystal. The development of the fiber structure under high drawing stress was almost complete in less than 1 ms, and the developed structure contained only β -crystals. Small-angle X-ray scattering images showed meridional streaks at low drawing stress, whereas a four-pointed pattern occurred under high drawing stress. Low drawing stress generated a long periodic structure that was defective in the periodic regularity of crystalline and amorphous regions, although the molecular chains were nearly oriented along the fiber axis. The high drawing stress resulted in a well-packed structure of adjacent fibrils with alternating amorphous and crystalline regions.

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INTRODUCTION

Poly(vinylidene fluoride) (PVDF) is a noteworthy engineering plastic owing to its excellent properties; namely, good chemical resistance, high mechanical strength and abrasion resistance. In particular, PVDF fiber can be melt-spun to form a fishing line because of its low water absorption, low refractive index similar to that of water, high specific gravity and good resistance to degradation by light. PVDF has been identified to take on five crystalline forms: α , β , γ , δ and ϵ .^{1–4} The α form (form II) is formed by cooling a quiescent melt, and it forms a nonpolar pseudo-orthorhombic structure containing two chains of tg^+tg^- conformation. Mechanical drawing of the α -form produces the β form (form I) containing two all-*trans* chains. Other forms can be produced by the application of electrical fields.^{5,6}

Some studies on the development of structure in PVDF fibers and tapes during melt spinning and deformation have been conducted using simultaneous synchrotron radiation as well as X-ray from laboratory generators for wide-angle X-ray diffraction (WAXD) and small-angle X-ray scattering (SAXS).^{7–10} Cakmak and co-workers⁷ have shown that the appearance of the β -phase is stress dependent, first occurring at a spinline stress of 3 MPa and increasing in concentration as stress is increased. The superstructure of the fibers was investigated using small-angle light scattering, where a transition from a spherulitic structure to a sheaf and rod-like

structure was observed with increasing draw ratio. In a later study, Cakmak *et al.*⁸ suggested that PVDF crystallizes with a shish-kebab-type morphology, which is a model consisting of a combination of an extended chain crystal (a shish) and folded chain crystals (kebabs),¹¹ during the melt-spinning process. Schultz and co-workers¹⁰ analyzed the crystal transformation from the α - to the β -form and morphological changes in lamellar and fibrillar structures during deformation at room temperature, and they found that deformation assisted in the conversion from the α - to the β -form. More α - to β -transformation took place in necked regions than in unnecked regions.

Our research group has studied the development of the fiber structure in laser-heated drawing for PET running fibers through WAXD and SAXS analyses¹² and afterward for PET and PTT fibers using *in situ* synchrotron WAXD and SAXS measurement systems.^{13,14} The *in situ* measurements were conducted as a function of elapsed time calculated from the distance between the measurement and necking positions with high time resolution because the necking could be fixed within a highly confined region by rapid and uniform heating with CO₂ laser radiation (see Figure 3). We succeeded in conducting *in situ* analyses with time resolutions of 1 ms for X-ray diffraction characterization and 0.47 ms for fiber temperature measurements in PET fibers. This study investigated the development

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of fiber structure in the vicinity of the necking location for PVDF fiber using our laser-drawing system and *in situ* measurement techniques.

EXPERIMENTAL PROCEDURE

Fundamentals of *in situ* measurement

Figure 1 shows a schematic diagram of the *in situ* measurement system used in this study. Details of the *in situ* system can be found in our previous papers.^{12–14} The running fiber was heated by irradiation with a CO₂ laser beam generated by a PIN-20S laser source manufactured by Onizca Glass (Tokyo, Japan). This laser source has a rated power of 20 ± 1 W and a laser beam diameter of 5 mm. The running fiber was drawn by speed difference between the feed and take-up rollers. The position of necking induced by laser irradiation was controlled within 1 mm during the drawing. X-ray diffraction images were recorded, and fiber temperature was measured as a function of elapsed time t , which was calculated from the distance D between the necking and measurement positions divided by the fiber running speed v . The distance D varied with necking position, which was changed by moving the laser irradiation.

The time resolution of the continuous drawing was calculated to attain a confidence interval at every measurement point. The time resolution was obtained by dividing the position resolution by take-up speed (m min^{-1}). The position resolution $W_{\text{X-ray}}$ can be calculated using equation (1) based on the width of the X-ray beam, W_{beam} , the fluctuation of the necking point, W_{neck} , and the length of the necking range, W_{deform} .

$$W_{\text{X-ray}} = \sqrt{W_{\text{beam}}^2 + W_{\text{neck}}^2 + W_{\text{deform}}^2} \quad (1)$$

In this case, W_{beam} and W_{deform} were 340 and 100 μm , and W_{neck} ranged from 100 to 300 μm . The resulting time resolutions for low and high drawing stress were 0.69 and 0.28 ms, respectively.

Materials and drawing conditions

The PVDF fiber of 250 μm diameter was kindly provided by Kureha Gohsen (Tochigi, Japan) and drawn to draw ratios of 4.5 and 5.5 with corresponding take-up speeds of 27 and 66 m min^{-1} , respectively. Table 1 shows the drawing conditions used in this experiment. Drawing stress was measured by a three-points-type tensiometer (HS-1500S; EIKO SOKKI, Osaka, Japan), and the fiber was drawn under two different drawing stresses: that is, high and low drawing stresses of 242 and 110 MPa, respectively. The applied laser

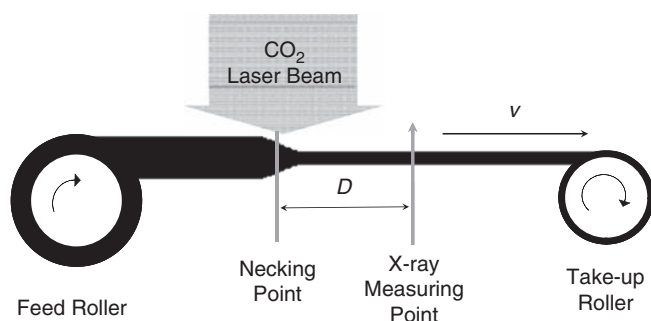


Figure 1 Schematic diagram of the on-line measurement system. The elapsed time after the neck deformation can be calculated from the distance D between the neck deformation location and the measuring positions divided by the fiber running speed v .

Table 1 Drawing conditions for the laser drawing of PVDF fibers

Drawing stress (MPa)	Draw ratio	Feed speed (m min^{-1})	Take-up speed (m min^{-1})	Time accuracy (ms)
110	4.5	6	27	0.69
242	5.5	12	66	0.28

Abbreviation: PVDF, poly(vinylidene fluoride).

intensity was measured using a power meter (PW-250; SYNRAD, Mukilteo, WA, USA).

Fiber temperature measurement

Fiber temperature was measured using an infrared thermo-spot-sensor (TMZ7-J0-2.5K0.2 type; Japan Sensor, Tokyo, Japan) equipped with a mercury-cadminto-telluride (MCT) detector and an interference filter. The spot size was 200 μm in diameter, and the response time was 10 ms. The interference filter transmits 3–4 μm in wavelength, corresponding to an absorbance band (2500–3333 cm^{-1}) representing C–H stretching. The fiber temperature was calculated from the radiation intensity averaged over 30 s for each measurement point. The fiber temperature was calculated in the same way as in the previous study.^{12–16}

X-ray diffraction characterization

X-ray diffraction measurements were carried out using the synchrotron radiation of BL40B2 in SPring-8 (Hyogo, Japan). The applied beam was 0.1 nm in wavelength and 340 μm in width. Diffraction images were taken using a 3000 \times 3000 pixel imaging plate attached to a 400-mm-long vacuum chamber. The sample-to-detector distance of 419.8 mm was estimated using lead stearate. The exposure time was 4 min. The neck deformation position was determined by a transmitted X-ray intensity profile through an ion chamber. X-ray images containing diffractions of polyimide film mounted on the vacuum chamber were compensated for by subtracting a blank image.

RESULTS AND DISCUSSION

Neck deformation behavior

Figure 2 shows images from CCD videos taken in the vicinity of the neck deformation. These videos revealed two distinctive appearances depending on drawing stress, with a moderate neck deformation under a low drawing stress of 110 MPa and a steep neck deformation under a high drawing stress of 242 MPa.

CO₂ laser absorption coefficient

The CO₂ laser absorption coefficient for PVDF was characterized using an infrared spectrometer (Shimadzu FT-IR 8400, Shimadzu,

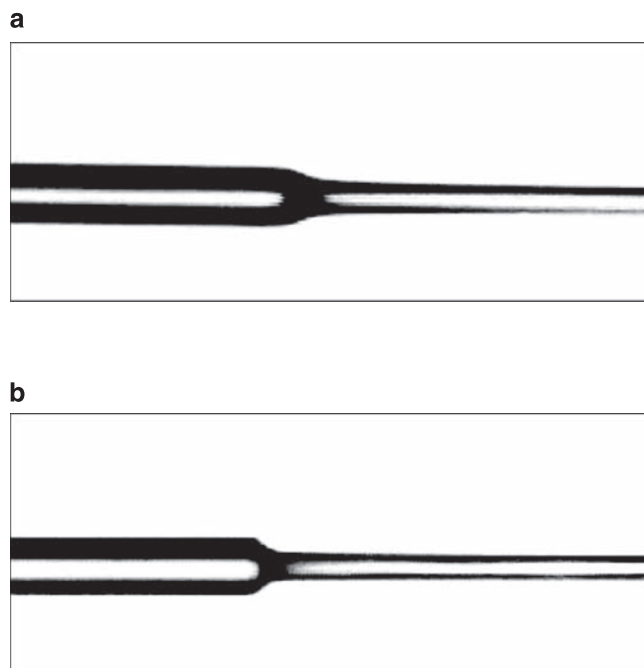


Figure 2 CCD video images in the vicinity of neck deformation with low (a) and high (b) drawing stresses.

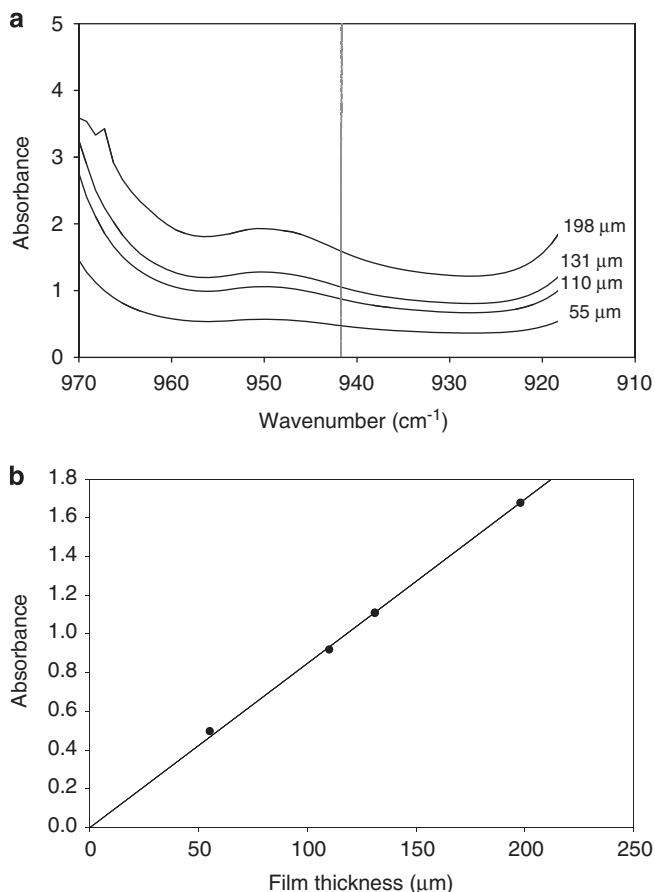


Figure 3 Infrared absorption spectra (a) and absorbance at 943 cm^{-1} (b) as a function of film thickness. Film thicknesses are noted on the right side of individual spectra.

Kyoto, Japan). The CO_2 laser used in this study was $10.6\ \mu\text{m}$ in wavelength, corresponding to an absorbance of $943\ \text{cm}^{-1}$. Infrared absorption spectra were obtained for four PVDF films that were 55, 110, 131 and $198\ \mu\text{m}$ in thickness, and the absorbance at $943\ \text{cm}^{-1}$ was extrapolated based on film thicknesses. The results are shown in Figures 3a and b. The infrared spectra were obtained by 10 integrations with a resolution of $2\ \text{cm}^{-1}$. Ultimately, the absorption coefficient for PVDF was found to be $1.908 \times 10^4\ \text{m}^{-1}$.

Fiber temperature profile

Figure 4 shows the fiber temperature profile over time in continuous laser drawing with a high drawing stress at a draw ratio of 5.5, where 0 on the horizontal axis indicates the time the necking occurred and the gray-shaded area denotes the region of laser irradiation. The fiber temperature was measured twice to reduce the experimental error, and this figure shows these two results together. Before the necking, the fiber temperature increased slightly due to the laser irradiation. At the necking point, that is, at an elapsed time of 0 ms, the fiber temperature showed a sharp increase from about 65 to 85°C , which may be the result of the plastic deformation of the necking. The fiber temperature continued to increase until 2 ms after necking, when the fiber was leaving the laser irradiation area. The exothermic crystallization seems to have finished at that point. Although the PVDF has a negative T_g of about -35°C , the necking occurred in the range of 65 – 85°C because it is a crystalline polymer. The orientation in the crystalline and

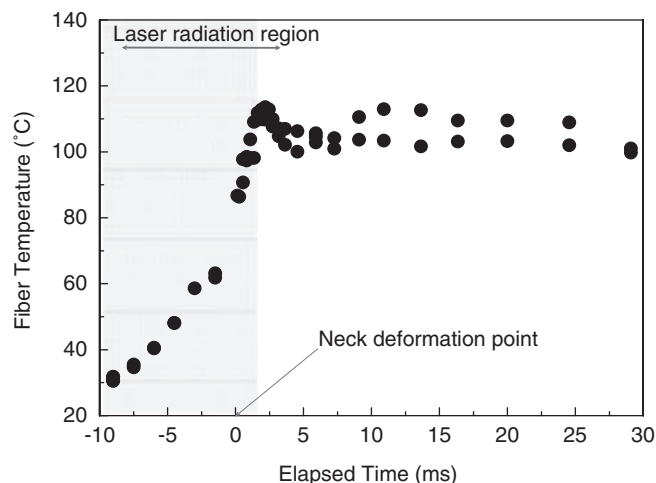


Figure 4 Fiber temperature profile as a function of elapsed time, where 0 on the horizontal axis indicates neck deformation and the gray-shaded area denotes the laser-irradiated region.

amorphous regions increased along the fiber axis with neck deformation. After an elapsed time of around 3 ms the fiber temperature was almost saturated, indicating that the development of the fiber structure was complete.

X-ray diffraction

WAXD analyses. Figures 5 and 6 show X-ray diffraction images of PVDF fibers drawn with low and high drawing stresses, respectively, and their X-ray intensity profiles are shown in Figures 7 and 8, respectively. The X-ray diffraction images revealed a typical PVDF crystal with α -(100)/(020), α -(110), β -(110)/(200) and α -(021) planes. Before the necking, the WAXD images showed an unoriented α -crystal. In Figure 5, the α -(110), β -(110)/(200) diffractions began to concentrate around the equator just after the neck deformation, and the α -(100)/(020) diffraction was concentrated on the equator at 0.5 ms. This result seems to indicate that the diffractions of α -(110), β -(110)/(200) and α -(100)/(020) began to develop immediately after the neck deformation. The α -(021) plane was oriented on the off-meridian after 2.0 ms, which could result from the tg^+tg^- conformation of the α -crystal. Figure 7 contains three diffraction peaks along the equatorial direction corresponding to the α -(021), α -(100)/(020), α -(110) and β -(110)/(200) diffractions. The α -(021) diffraction disappeared with neck deformation at the equator. The α -(110) and β -(110)/(200) planes were close to each other and were located in only one peak. Just after the neck deformation, the α -(110), β -(110)/(200) diffraction became somewhat broad and shifted to a higher 2θ . This shift was caused by the fact that the β -(110)/(200) diffraction intensity increased with elapsed time. Subsequently, the β -crystal was developed immediately after the neck deformation and increased with elapsed time. Ultimately, the β -crystal was dominant in the drawn fiber, although the α - and β -crystals coexisted in the laser-drawn PVDF fiber with a low drawing stress of 110 MPa. Nagaoka *et al.*¹⁷ have reported that the development of crystal structure in laser-drawn PVDF fibers depends on the drawing stress and that drawing stresses higher than 100 MPa largely produce β -crystals. Therefore, the β -crystals developed earlier than the α -crystals in the neck drawing: the β -crystal developed within 1 ms after the neck deformation. In Figure 6, the α -(110) and β -(110)/(200) diffractions are concentrated on the equator at 0 ms and are well separated from the adjacent

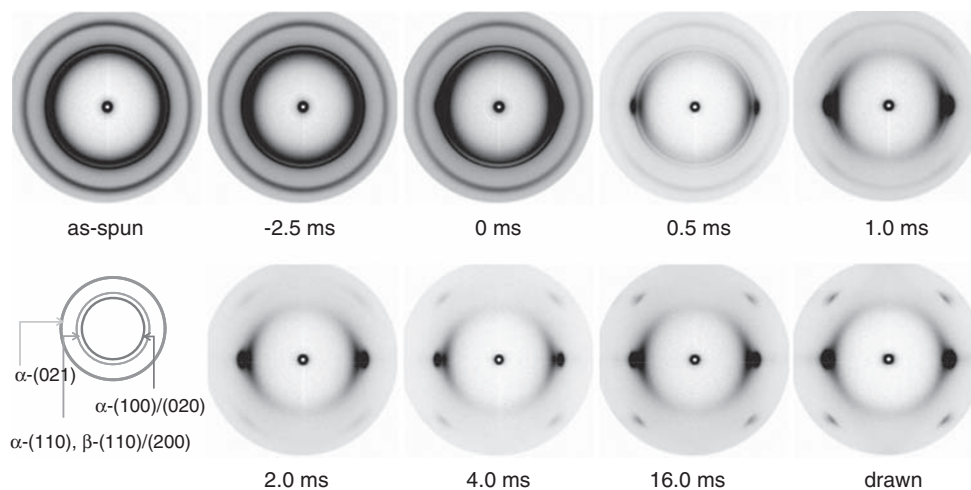


Figure 5 X-ray diffraction images taken at various elapsed times during continuous laser drawing under low drawing stress.

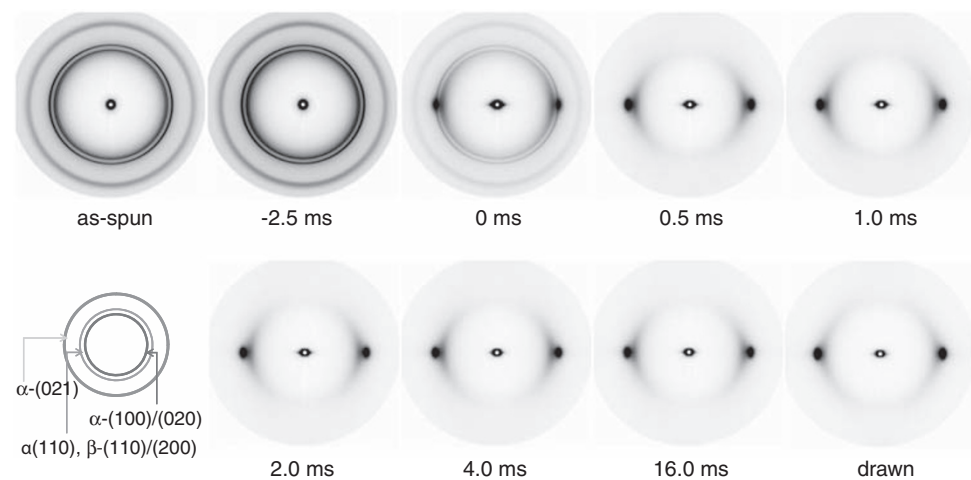


Figure 6 X-ray diffraction images taken at various elapsed times during continuous laser drawing under high drawing stress.

diffraction, unlike the low drawing stress condition. The diffraction pattern at 0.4 ms clearly reveals the β -crystal formation, and then the pattern hardly changes with elapsed time. Accordingly, the fiber structure development was complete immediately after the neck deformation. The α -crystal diffractions were weak at and after 0.4 ms, and they faded out in the drawn fiber. In Figure 8, we can observe only β -crystals in the drawn fiber.

SAXS analyses. The X-ray diffraction patterns showed some changes in both the SAXS and the WAXD patterns because this characterization used a long distance path of 419 mm. Therefore, we were able to successfully collect SAXS patterns by magnifying the center portion of the X-ray patterns, from which qualitative SAXS analyses can be conducted. Figures 9 and 10 show SAXS images taken in laser drawing with low and high drawing stresses, respectively, and these images show the large-scale periodic character of the structure parallel to the fiber axis. The as-spun fiber showed an appearance of curvature at the meridian, indicating a substantial misorientation of lamella with respect to the fiber axis. In Figure 9, the curvature decreased at 0 and 0.5 ms and then increased and flattened off with elapsed time.

Ultimately, the drawn fiber had an appearance of meridional streaks, which may be a precursor of a four-pointed scattering pattern. Thus, the molecular chains were oriented nearly parallel to the fiber axis, and crystalline and amorphous regions alternate in a rather regular manner along the fibril axis. Therefore, in the laser drawing with a low drawing stress of 110 MPa, the long period structure is somewhat defective in the periodic regularity of crystalline and amorphous regions. Adjacent fibrils in the low-stress condition seemed not to be as close as those in the high-stress condition, and the fibers generated under low stress seemed to be generated after 1 ms.

In Figure 10, on the other hand, the curvature disappeared at the same time as the initiation of neck deformation. SAXS patterns at and after 0 ms presented equatorial streaks, which can be attributed to microvoids generated by drawing under high stress.¹⁰ The streaks decreased with elapsed time, indicating that the microvoids were reduced and concentrated. At and after 0.5 ms, a four-pointed pattern was observed and it became increasingly clear with elapsed time. This pattern occurs because adjacent fibrils packed together by alternating amorphous and crystalline regions immediately after the neck deformation, and these regions then became fastened together as time

elapsed. In the drawn fiber, the equatorial streak existed clearly even though the four-pointed pattern was completed, and thus, the high drawing stress of 242 MPa may result in a large quantity of microvoids among interlocking fibrils. Furthermore, the high drawing stress may transform the initial lamellar structure into a fibrillar structure after 0.5 ms. This result indicates that the development of the fibrillar structure of PVDF fibers depended on both the drawing stress and the elapsed time.

Conclusion

The development of fiber structure in the vicinity of the necking location of PVDF fibers was investigated as a function of elapsed time using the laser-drawing system and *in situ* measurement techniques.

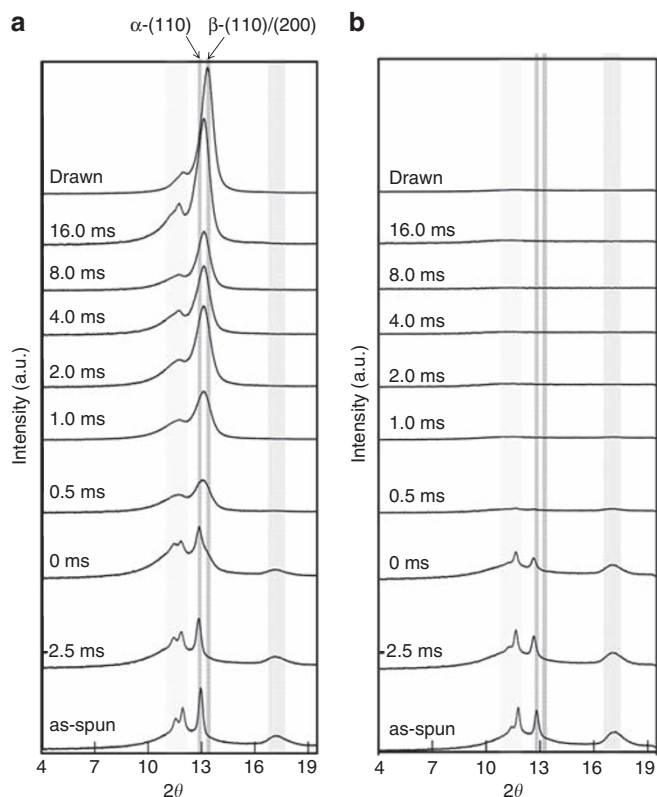


Figure 7 X-ray intensity profiles along the equatorial (a) and meridional (b) directions for low drawing stress.

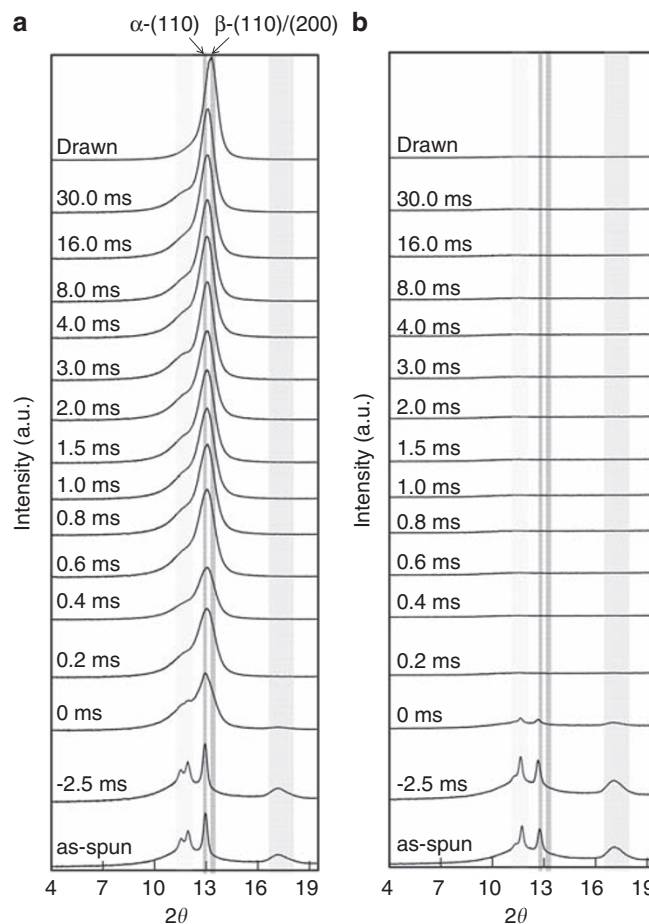


Figure 8 X-ray intensity profiles along the equatorial (a) and meridional (b) directions for high drawing stress.

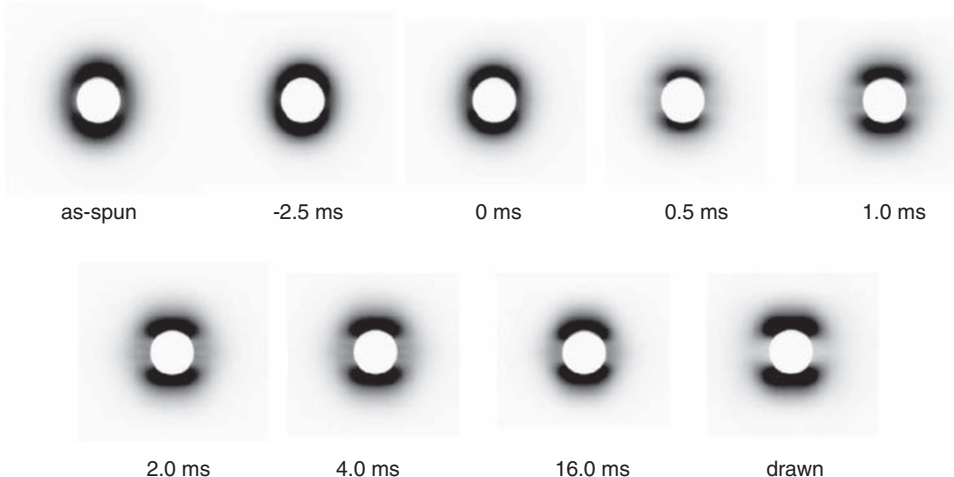


Figure 9 SAXS patterns for low drawing stress (magnified images of Figure 6).

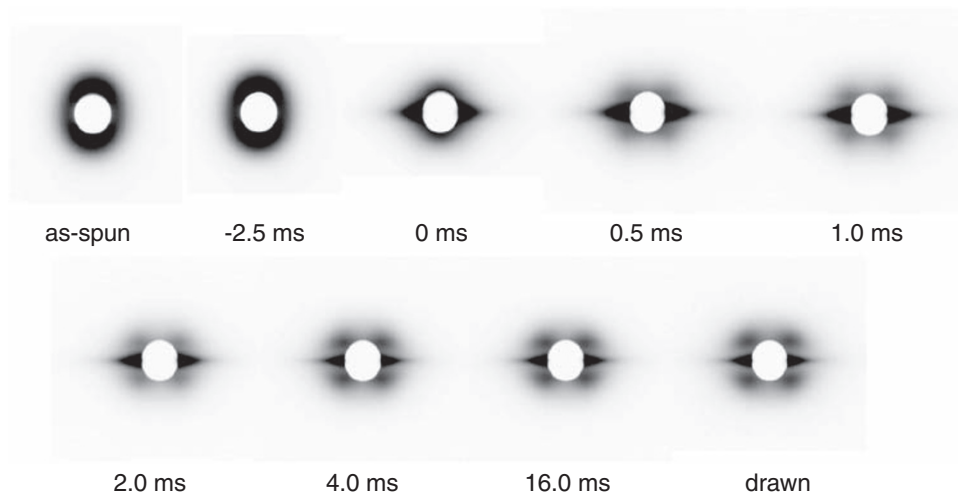


Figure 10 SAXS patterns for high drawing stress (magnified images of Figure 7).

As the necking position was fixed within a highly confined region by rapid and uniform heating with CO₂ laser radiation, we succeeded in conducting *in situ* analyses with time resolutions of 0.69 and 0.28 ms in the laser-heated drawing of PVDF fibers for drawing stresses of 110 and 242 MPa, respectively. The development of the crystal structure observed in WAXD analysis showed that the β -crystal developed immediately after the neck deformation and continued to develop with elapsed time. Ultimately, β -crystals were predominant in the drawn fiber, although the α - and β -crystal forms coexisted in the laser-drawn PVDF fiber produced with a low drawing stress of 110 MPa. For the laser-heated neck drawing of PVDF fibers with a high drawing stress of 242 MPa, the fiber structure development was complete immediately after the neck deformation, and only β -crystals were observed in the drawn fiber. The supramolecular structure elucidated by SAXS indicated that the laser drawing of PVDF fibers with a low drawing stress resulted in a defective long period structure in which adjacent fibrils were farther apart from each other than in the high-stress condition. Laser drawing with a high drawing stress produced a four-pointed pattern with an equatorial streak immediately after the neck deformation; that is, the high drawing stress produced a long period structure with a large quantity of microvoids among interlocking fibrils. The fiber structure development in the laser drawing of the PVDF fiber depended on the drawing stress and on the elapsed time after necking.

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