

Supporting Information

Deformation Mechanism of Poly(3-alkylthiophene) Studied by *in Situ* X-ray

Scattering and Texture Analysis

Haiming Chen,^{†,‡} Cui Su,^{†,‡} Guangyu Shi,^{†,‡} Xiaolu Li,[§] Xiuqin Zhang,[§] Guoming Liu,^{*,†} and
Dujin Wang^{†,‡}

[†] CAS Key Laboratory of Engineering Plastics, CAS Research/Education Center for Excellence in Molecular Sciences, Institute of Chemistry, Chinese Academy of Sciences, Beijing 100190, China

[‡] University of Chinese Academy of Sciences, Beijing 100049, China

[§] Beijing Key Laboratory of Clothing Materials R & D and Assessment, Beijing Engineering Research Center of Textile Nanofiber, School of Materials Science & Engineering, Beijing Institute of Fashion Technology, Beijing 100029, China

Corresponding author: gmliu@iccas.ac.cn

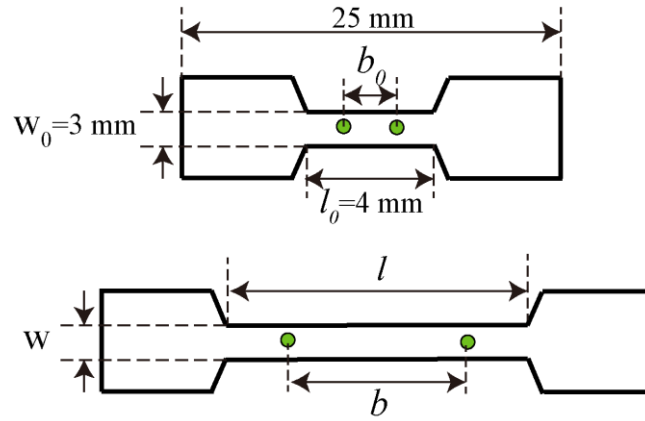


Figure S1. The dimension and marker points (green circles) on the surface of mini tensile bar before and after stretching.

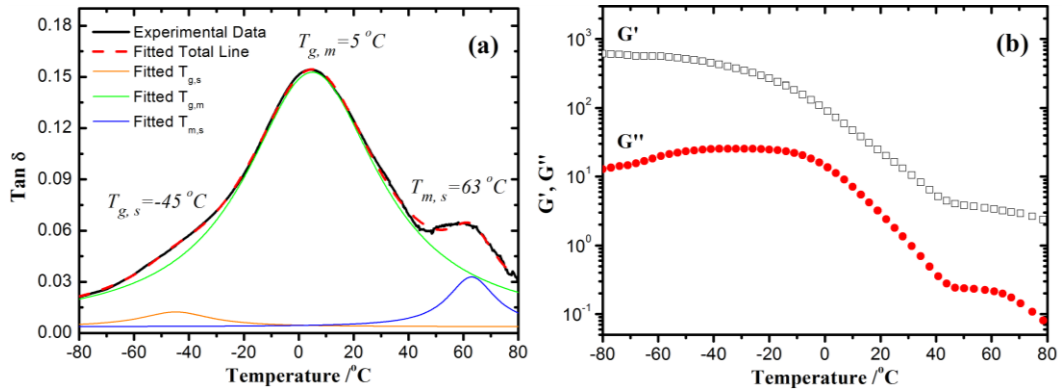


Figure S2. The (a) $\tan \delta$ and (b) storage modulus (G'), loss storage (G'') of P3DDT film obtained from dynamic mechanical analysis (DMA) measurement. DMA was carried out with a TA Q800 instrument employing uniaxial tension mode. The temperature scan rate was 3 °C/min and the scan range was -100 °C ~ 100 °C. The dimension of the specimen was 11.90 mm \times 5.92 mm \times 0.06 mm (length \times width \times thickness). The frequency was fixed at 1 Hz. A strain of 0.1 % was chosen which was located in the linear viscoelastic regime.

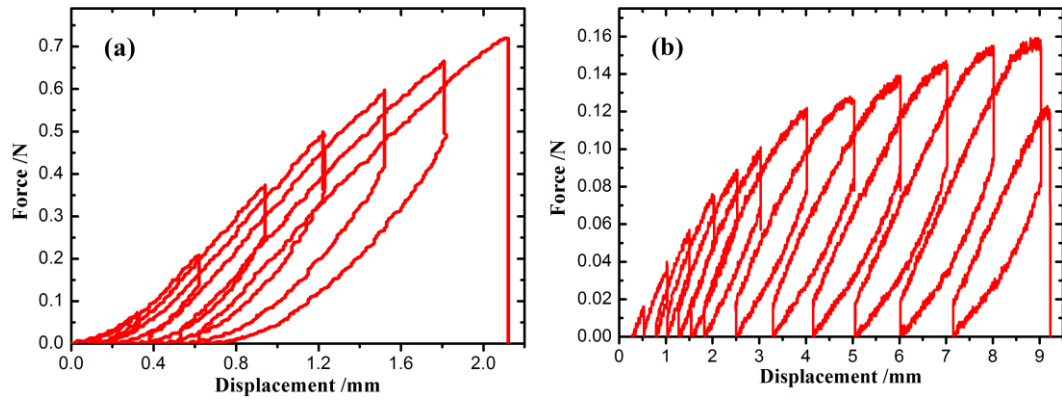


Figure S3. Force-displacement curves of the step-cycle measurements of P3DDT at 30 °C (a) and 110 °C (b), respectively.

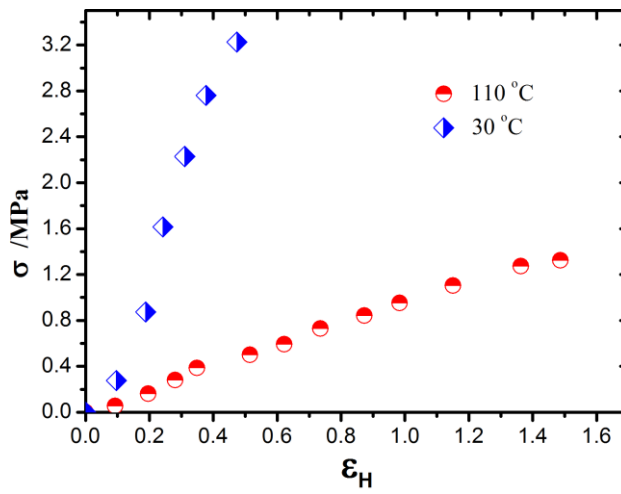


Figure S4. True stress (σ)-strain (ϵ_H) plots of P3DDT film at 30 °C and 110 °C, respectively.

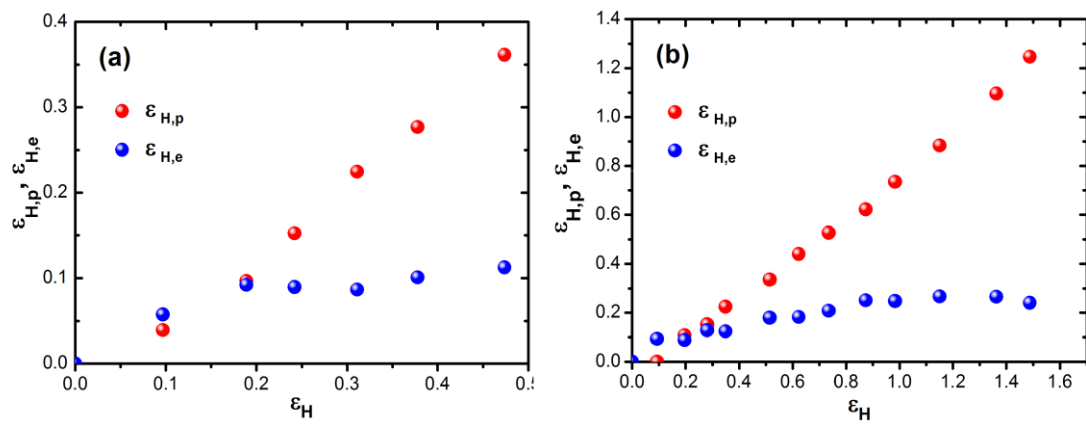


Figure S5. Elastic ($\epsilon_{H,e}$) and plastic ($\epsilon_{H,p}$) strain as a function of total strain (ϵ_H) of P3DDT at (a) 30 °C and (b) 110 °C, respectively.

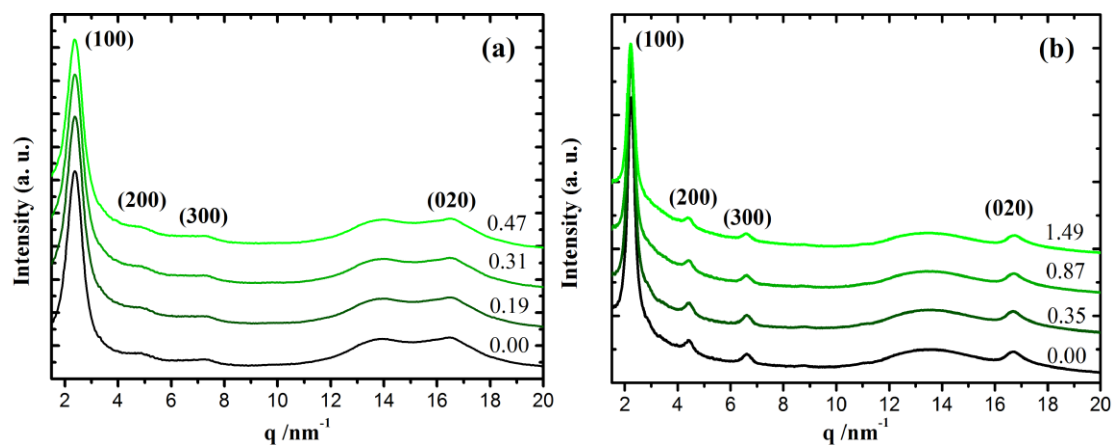


Figure S6. 1D WAXS intensity profiles of P3DDT upon stretching at 30 °C (a) and 110 °C (b), respectively.

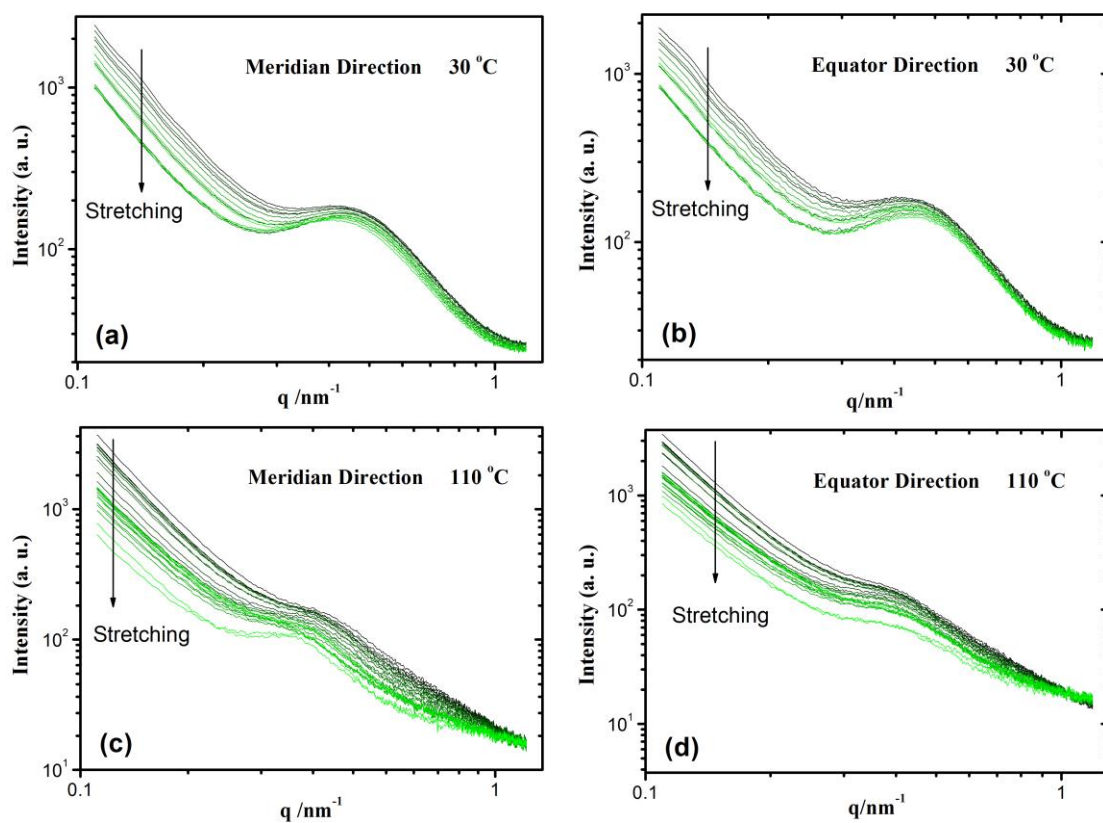


Figure S7. 1D SAXS intensity profiles of P3DDT under uniaxial stretching at 30 °C (a)-(b) and 110 °C (c)-(d).

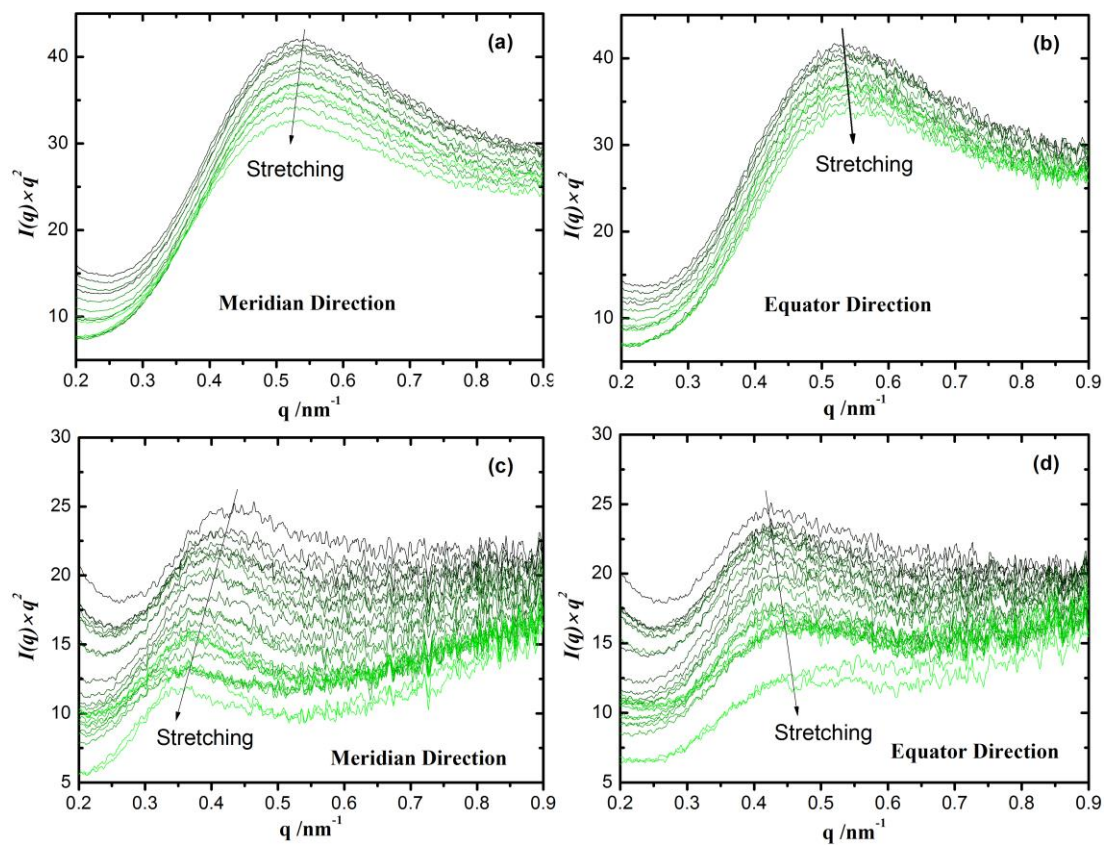


Figure S8. Lorentz corrected SAXS profiles of P3DDT under uniaxial stretching at 30 °C (a)-(b) and 110 °C (c)-(d).

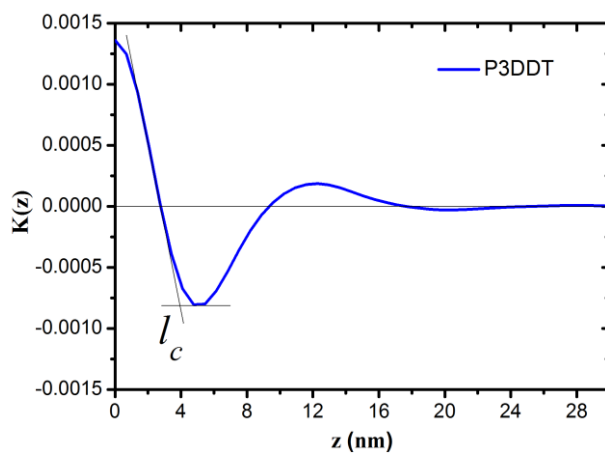


Figure S9. A typical one-dimensional electron density correlation function of unstretched P3DDT at 30 °C. l_c is crystalline layer thickness, which can be estimated from the correlation function curve.

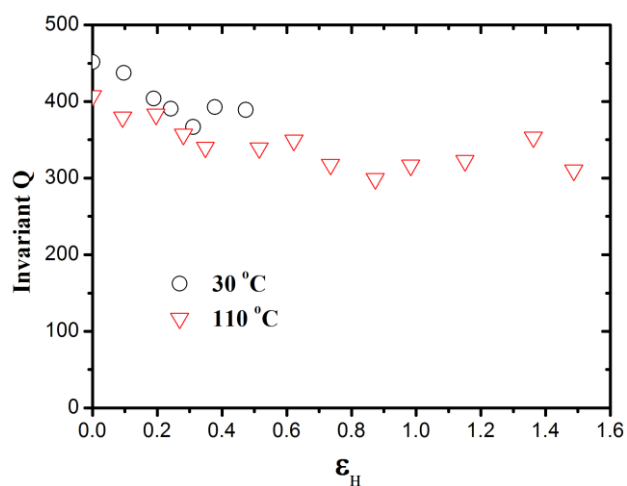


Figure S10. Scattering Invariant Q as a function of true strain (ϵ_H) at different temperatures.

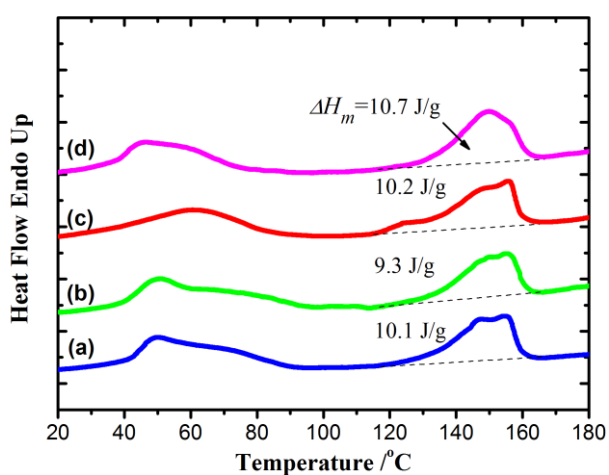


Figure S11. DSC curves of P3DDT before and after stretching. (a) and (c) are annealed at 30 °C and 110 °C for 30 min before stretching, respectively. (b) and (d) are after stretching at 30 °C and 110 °C, respectively. DSC measurements were carried out on a Q2000 DSC (TA Instruments). Temperature scans were performed within the range of 0 °C to 200 °C. Around 4 mg sample were used for measurement. The instrument was calibrated with indium.