Supporting Information

Structural Evolution Analysis and Cold-Crystallization Kinetics to Spherical Crystal

in Poly (trimethylene terephthalate) Film using Raman Spectroscopy

Chenglong Hu,^{a,b,} Shaoyun Chen, *aWeihong Zhang, Fangyan Xie, Jian Chen and Xudong Chen,^b

^{*a*}Key Laboratory of Optoelectronic Chemical Materials and Devices of Ministry of Education, Jianghan University, Wuhan 430056, China.

^{*b*}Key Laboratory for Polymeric Composite and Functional Materials of Ministry of Education, DSAPM Lab, School of Chemistry and Chemical Engineering, Sun Yat-sen University, Guangzhou 510275, China. ^{*c*}Instrumental Analysis and Research Center, Sun Yat-sen University, Guangzhou 510275, China.

*Corresponding authors: cescsy@163.com (S. Y. Chen); Key Laboratory of Optoelectronic Chemical Materials and Devices of Ministry of Education, Jianghan University.



Figure S1 (a) and (b) Typical FTIR spectra of an amorphous PTT film at certain temperatures. (c) FTIR absorbance of an amorphous PTT film at 2967 cm⁻¹ with a function of temperature (the heating rate was 5 °C/min in air). Cp and Cs represented primary crystallization and secondary crystallization, respectively. (d) DSC heating versus temperature of an amorphous PTT film collected at a heating rate of 5 °C/min in nitrogen).



Figure S2 The Raman spectra of an amorphous PTT film at certain isothermal crystallization temperature (T_c) (a) $T_c=68$ °C, (b) $T_c=70$ °C, (c) $T_c=74$ °C and (d) $T_c=78$ °C.



Figure S3 Dependence of relative crystallinity on crystallization time (solid line) and its differential curve (dash line). Crystallization temperature was 68 °C. The dynamic process contained four phases: (N) nucleation phase, (NG) nucleus growth phase and (Cs) secondary crystallization phase.