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Application of the correct design of successive self-nucleation and annealing (SSA) to study the stereo-defects and its distribution of homo- and co-polypropylene

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The stereo-defects and its distribution of four homo- and co-polypropylene samples with different processing properties were studied through the correct design Successive Self-Nucleation and Annealing (SSA), and other characterization methods such as Differential Scanning Calorimetry (DSC), Gel Permeation Chromatography (GPC), Temperature Rising Elution Fractionation (TREF), respectively. Firstly, the preliminary characterization found that the two homo-polypropylene or co-polypropylene samples had the similar mechanical and thermal properties, and the SSA results also revealed the internal microstructure of the different kind of polypropylene was almost exactly the same, representing the alike isotactic sequence length, the fractions weight of the isotactic sequence, and the isotactic sequence length distribution of the same kind of polypropylene. Compared with the homo-polypropylene, the DSC, GPC, and TREF tests indicated there were difference in the properties of the co-polypropylene samples, revealing the disparities in molecular chain microstructure. The SSA results after the correct T_g for each sample showed there were obvious differences in the (statistical) lamellar thickness and its distribution and the fractions weight of different fractions after SSA treatment, further revealing the microstructure difference on the isotactic sequence length, and the isotactic sequence length distribution of homo- and co-polypropylene samples. Besides, the correct design SSA also indicated the polar copolymer monomer had a great influence on the high isotactic component in co-polypropylene microstructure.

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Introduction

Polypropylene was usually prepared by Ziegler–Natta catalyst system and widely used in various applications such as packaging material and other industrial uses such as lithium-ion battery separators, and the stereo-defects of polypropylene could reflect the information the processability and mechanical properties of polypropylene.^{1–3} It is known that the methyl groups of its molecular chain for polypropylene are not all perfectly in the same side, and the stereo-defects of polypropylene prepared by Ziegler–Natta catalyst are the stereo-defects of the arrangements of the side methyl groups of “mistake inserted” monomers, and the length between every two “mistake inserted” monomers is called isotactic sequence length of polypropylene, as shown in Fig. 1, which can reflect stereo-defects distribution of polypropylene.^{4–10}

Successive Self-Nucleation and Annealing (SSA) was a thermal fractionation protocol, which was designed and implemented by Müller and coworkers in 1997.^{11–19} SSA was widely used to analyze the chain structures of crystallized polymers (*e.g.*, branches,

comonomers, crosslinks, stereo-defects or any other molecular defects) based on the self-nucleation and annealing steps, which was capable of performing thermal fractionation with much better resolution and spending less time than previously available techniques (such as step crystallization and TREF). And there was a good correspondence between the SSA and TREF techniques, and the results of SSA were coincident with the results of high resolution ¹³C-NMR, GPC, TREF, and xylene solvent (XS) fractionation.^{13,16–34}

For PP samples, with the increase of separation temperature, the component separated by TREF showed higher melting temperature and longer isotactic sequence length after SSA treatment, and ¹³C-NMR result also indicated that the isotacticity of polypropylene and isotactic sequence length of PP increased.^{27,35–38} Compared with ¹³C-NMR and TREF fractionation, SSA fractionation was a more time-saving and practical technique

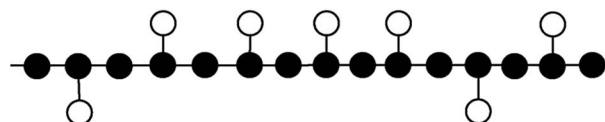


Fig. 1 The schematic diagrams of PP chain structure.

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in the characterization of the molecular structure of PP, as well as a way providing important quantitative information about the lamellar thickness and its distribution of PP, which gave a more accurate characterization to the isotactic sequence length and its distribution of polypropylene.

However, Müller found that some authors started their SSA protocol at an arbitrary T_s without a previous self-nucleation study to determine the ideal self-nucleation temperature ($T_{s\text{-idea}}$) although there were more than 130 reports on the application of SSA on the polymers since 1997, which was reported as the review on the principles, applications and perspectives of SSA by Müller.³⁹ Müller insisted that any attempts to quantify the data such as calculate SCB or MSL distributions were not correct unless the first T_s employed to perform SSA was $T_{s\text{-idea}}$ because the $T_{s\text{-idea}}$ had a strong impact on the final fractionation profile. Therefore, correct design of the fractionation method should include a conscientious choice of fractionation window, fractionation time, scanning rate, sample mass and especially the first self-nucleation temperature (T_s) to be employed, and the self-nucleation studies must be performed before SSA in order to choose the first T_s .

In this paper, the stereo-defects and its distribution of four homo- and co-polypropylene samples with different processing properties were studied through the correct design of Successive Self-Nucleation and Annealing (SSA) and other ways such as DSC, GPC, TREF, respectively. For SSA fractionation profile, the fractionation window was 5 °C, the fractionation time was 15 min, the scanning rate was 20 °C min⁻¹, and sample mass was 2.5 mg according to the high-speed calorimetry concept introduced by Pijpers.^{11,40–42} So, correct design of the fractionation method for the first self-nucleation temperature (T_s) was employed by self-nucleation studies (SN) before SSA in order to choose the first T_s .

Experimental

Materials

The four homo- and co-polypropylene samples studied in this paper were commercial PP resin. The homo-polypropylene sample 1 (grade T30S) and the co-polypropylene sample 3 (grade k8003) were provided by China Sinopec, while the sample 2 (grade PC366-3) and the co-polypropylene sample 4 (grade 7633) were provided by LCY Chemical Corp., respectively.

Characterization of the four homo- and co-polypropylene samples

The molecular weights (M_n and M_w) and the molecular weight distribution (MWD) were determined by a PL-GPC 220 high-temperature gel permeation chromatography (Polymer Laboratories Ltd) at 413.15 K, using 1,2,4-trichlorobenzene as solvent, the injection volume was 100 mL, and the flow rate was 1.0 mL min⁻¹. Calibration was made by polystyrene as the standard sample.

The TREF tests of the PP samples were tested by a Polymer-Char TREF300, using 1,2,4-trichlorobenzene as solvent, and detection method is IR. The temperature was raised from 35 °C to 150 °C.

All differential scanning calorimetry (DSC) measurements were performed with a Mettler 822e DSC in a nitrogen atmosphere. Samples about 2–4 mg in aluminium pans were first melted by heating from 50 °C up to 200 °C at a rate of 50 °C min⁻¹ and were held at this temperature for 5 min in order to erase the previous thermal history, then cooled down to 50 °C at a rate of 20 °C min⁻¹, held at 50 °C for 5 min, and then finally heated again to 200 °C at a rate of 20 °C min⁻¹. The melting temperature T_m and the enthalpy of fusion ΔH_m were determined from the second melting curve. The degree of crystallinity was calculated according to the following formula (1):⁴³

$$\chi_c = \Delta H_m / \Delta H_m^0 \quad (1)$$

where ΔH_m was the fusion heat obtained from DSC curve, ΔH_m^0 was the fusion heat of a perfectly crystalline PP, *i.e.*, 209.0 J g⁻¹.⁴⁴

The self-nucleation (SN) technique was originally devised by Keller *et al.*⁴⁵ to aid the preparation of single crystals from solution. Fillon *et al.*⁴⁰ extended the study of self-nucleation by employing DSC. The SN procedures can be described as follows:

- Erasure of thermal history and crystalline memory. Erasure of the crystalline thermal history by heating the sample to 200 °C and held for 5 min;
- Creation of a “standard” semi-crystalline state. Cooling the sample at 20 °C min⁻¹ to 50 °C and held for 2 min;
- Thermal conditioning at a temperature denoted as T_s . The sample was heated at 20 °C min⁻¹ from 50 °C to a partial melting temperature denoted as T_s , and the sample was kept at T_s for 15 min;
- Subsequent cooling at 20 °C min⁻¹ from T_s down to 50 °C;
- Finally, the sample was heated at 20 °C min⁻¹ from 50 °C to 200 °C.

SSA fractionation was performed as presented by Müller.¹¹ Pijpers *et al.*⁴² recently introduced the high-speed calorimetry concept that the increment of heating rate can be compensated by reducing the sample mass. Hence, a higher scanning rate of 20 °C min⁻¹ was employed to reduce the fractionation time, and a smaller sample mass of 2.5 mg was used as compensation. The SSA procedure was schematically displayed in Fig. 2, and the complete thermal treatment comprised the following steps:

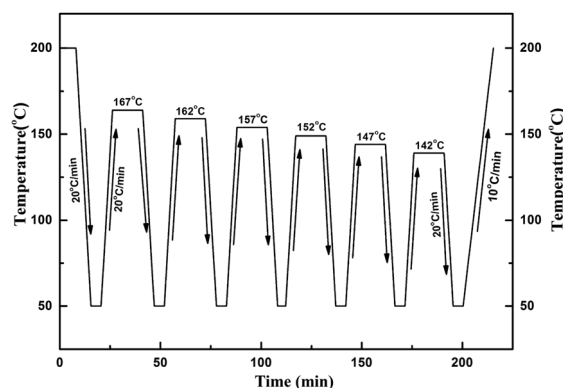


Fig. 2 The SSA thermal protocol schematic representation of sample 1.



- (a) Erasure of the crystalline thermal history by heating the sample to 200 °C and held for 5 min;
- (b) Cooling the sample at 20 °C min⁻¹ to 50 °C and held for 2 min;
- (c) The sample was heated at 20 °C min⁻¹ from 50 °C to a partial melting temperature denoted as T_s , such as the first T_s for sample 1 was chosen as 167 °C after the SN procedure;
- (d) The sample was kept at T_s for 15 min;
- (e) DSC cooling scan at 20 °C min⁻¹ from T_s to 50 °C;
- (f) Repeat step “c” to “e” at a new lower T_s which is 5 °C lower than the previous T_s for a total six self-nucleation/annealing steps.
- (g) Finally, the sample was heated at 10 °C min⁻¹ from 50 °C to 200 °C, and a multiple of melting endothermal peaks were obtained.

Results and discussion

The thermal properties characterization of PP samples

Firstly, all PP samples were characterized preliminarily and the results were listed in Table 1. The DSC melting curves and crystallization curves of four PP samples were shown as Fig. 3 and 4, respectively.

For the homo-polypropylene resin sample 1 and sample 2, Fig. 3, 4 and Table 1 indicated that there were subtle differences on the melt flow-ability, the melting temperature, the melting enthalpy and crystallization temperature of homo-polypropylene, revealing the microstructure difference of homo-polypropylene was small. Compared with sample 1, sample 2 had the higher melting temperature, the higher melting enthalpy, the higher crystallization temperature and degree of crystallization, suggesting that samples 2 had a regular molecular chain structure. For the co-polypropylene resin sample 3 and sample 4, Fig. 3, 4 and Table 1 revealed that the melting enthalpy, the crystallization temperature and the degree of crystallization of co-polypropylene were lower than those of homo-polypropylene, which was mainly ascribed to the regularity of the molecular chain of polypropylene destroyed by the copolymerization monomer.

The structure properties characterization of PP samples

The structure properties of all PP samples were studied by GPC and TREF, respectively.

Table 1 DSC characterization of four PP samples

Samples	MFR ^a	T_m^b (°C)	ΔH_m^c (J g ⁻¹)	T_c^d (°C)	X_c^e (%)
Sample 1	3.5	163.6	91.7	117.3	43.9
Sample 2	3.0	164.2	98.8	117.7	47.3
Sample 3	2.0	164.2	77.5	121.8	37.1
Sample 4	2.1	165.1	87.9	119.4	42.1

^a The MFR test condition of PP samples was g per 10 min, 2.16 kg, 230 °C. ^b Melting temperature was determined from the endothermic curves of DSC. ^c The value of the endothermic enthalpy was determined from DSC. ^d Crystallization temperature was determined from the exothermic curves of DSC. ^e The degree of crystallization was calculated from the value of the endothermic enthalpy.

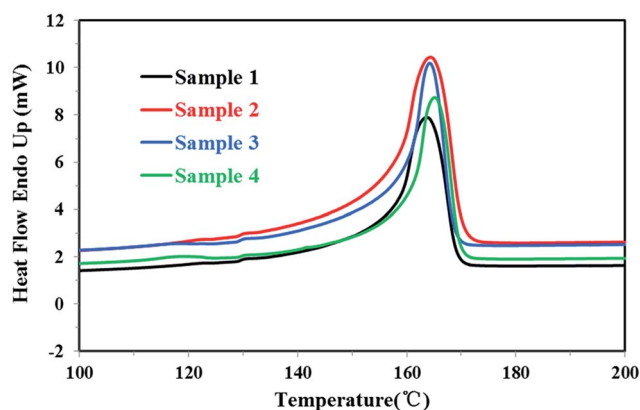


Fig. 3 The DSC melting curves of four PP samples.

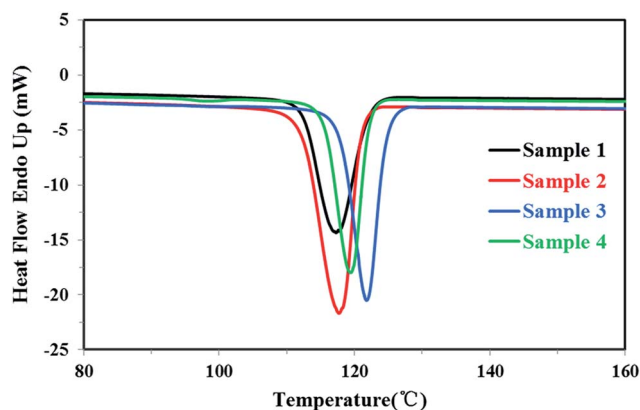


Fig. 4 The DSC crystallization curves of four PP samples.

The molecular weights (M_n and M_w) and the molecular weight distribution (MWD) of all PP samples were tested by GPC and plotted in Fig. 5 and Table 2. The GPC results revealed that the molecular weight distribution of homo-polypropylene was almost the same, which was about 2.7, but the molecular weights (M_n and M_w) of sample 2 was larger than that of sample 1, which may reveal there was an obvious difference on the regular molecular chain structure and molecular chain length between the homo-polypropylene samples 1 and samples 2. For

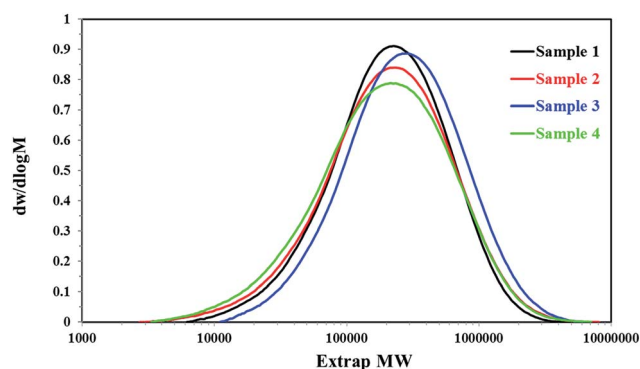


Fig. 5 The GPC trace of the different PP samples.



Table 2 GPC characterization of all PP samples

Samples	M_n	M_w	M_w/M_n
Sample 1	120 347	326 539	2.713
Sample 2	159 138	431 216	2.710
Sample 3	102 599	350 143	3.413
Sample 4	94 594	339 712	3.709

the co-polypropylene resin, sample 3 and sample 4, Fig. 5 and Table 2 showed that the molecular weight distribution of sample 3 was much smaller than that of sample 4, while the molecular weights (M_n and M_w) of sample 3 was heavier than that of sample 4, suggesting that the molecular chain of samples 3 was concentrated distribution on the chain segment of high molecular weight.

The TREF results of all PP sample were plotted in Fig. 6. The temperature was ranging from 40 °C to 140 °C. Compared with homo-polypropylene, there were obvious multiple melting endotherm (107.1 °C, 92.7 °C, 66.6 °C) of co-polypropylene respectively, indicating that there were more branched structures in co-polypropylene. As the TREF300 was an analysis TREF, which was not like the P-TREF that the fraction components could be collected and deeply characterized for microstructure characteristics, SSA fractionation was needed to give a more accurate characterization of the molecular structure of PP.

SSA thermal fractionation characterization of PP samples

As the ideal self-nucleation temperature (the first self-nucleation temperature T_s) had a strong impact on the final fractionation profile when the SSA thermal fractionation was used for polymers, for the correct design of the fractionation method, the self-nucleation studies must be performed before SSA in order to choose the first T_s .³⁹ According to Fillon's study,⁴⁰ the self-nucleation temperature range could be divided into three domains when the self-nucleation behavior of i-PP through the self-annealing procedures by DSC. Domain I can represent the temperature range where complete melting was achieved and the samples retained no memory of prior crystallization. Domain II was observed at lower T_s values, and the

nucleation density displayed an exponential growth and the peak crystallization temperature showed a corresponding shift to higher values when cooling from T_s . Domain III appeared at lower T_s temperatures where both partial melting and annealing occurred. Domain II applied to T_s temperatures which were high enough to melt almost any crystals in the sample, but still low enough to leave small crystal self-nuclei or seeds that could nucleate the polymer upon subsequent cooling from T_s . Müller *et al.*^{11,39} suggested that the optimum self-nucleation temperature or optimum T_s temperature was defined as the minimum temperature within Domain II of the SSA thermal conditioning.

According to standard self-nucleation procedure,^{11,39,40} the melting and crystallization curves of SN at different annealing temperature of sample 1 for example were studied, as shown in Fig. 7 and 8. Fig. 8 showed that there were no changes detected in the crystallization temperature obtained at the T_s temperature from 170 °C to 200 °C as compared to the standard crystallization temperature obtained at the T_s temperature of 200 °C, indicating that the nucleation density of the samples remained constant in Domain I. Thermal conditioning at 169 °C for 15 min was able to make polypropylene self-nucleation, so its crystallization temperature was shifted to higher temperature (Fig. 8) while the subsequent melting didn't reveal any traces of annealing (Fig. 7). That's because the unmelted crystal could form the self-nucleation of pp in the subsequent crystallization process, leading to significant reduction of the energy barrier of crystallization process. Fig. 7 showed that a new melting peak occurred in 174.1 °C when T_s was equal to and less than 166 °C, showing a typical single self-nucleation and annealing, which indicated that Domain III started. When the annealing temperature T_s was lower than 167 °C, the crystallization peaks became broader and the lamellae started to grow thicker. The lamellae were more perfect than the lamellae formed from the next dynamic crystallization process and had the higher melting temperature. The nucleation behavior of sample 1 illustrated that the minimum annealing temperature of Domain II was T_s value equal to 167 °C. The optimal annealing temperature T_s of all PP samples were tested as above self-nucleation experiments and the results were listed in Table 3.

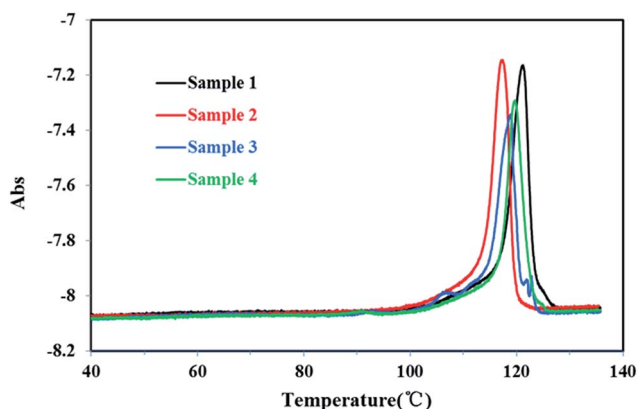


Fig. 6 The TREF trace of the different PP samples.

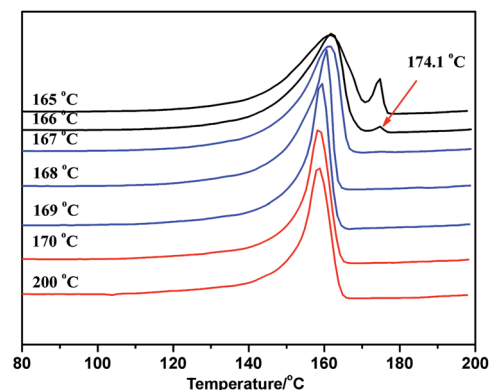


Fig. 7 The DSC melting curve after crystallization at different annealing temperature of sample 1.



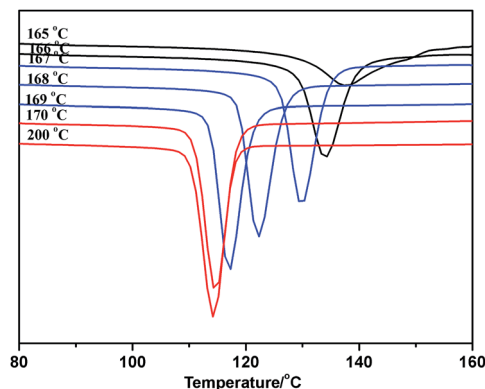


Fig. 8 The DSC crystallization curve after crystallization at different annealing temperature of sample 1.

Table 3 Domain-II and the optimal range of T_s of each sample

Samples	Domain-I	Domain-II	Domain-III	$T_s/^\circ\text{C}$
Sample 1	$T_s > 169$	$166 < T_s \leq 169$	$T_s \leq 166$	167
Sample 2	$T_s > 169$	$166 < T_s \leq 169$	$T_s \leq 166$	167
Sample 3	$T_s > 169$	$166 < T_s \leq 169$	$T_s \leq 166$	167
Sample 4	$T_s > 169$	$167 < T_s \leq 169$	$T_s \leq 167$	168

In order to study the distribution of stereo-defects in the polypropylene chain, all PP samples were studied and fractionated by SSA. The SSA melting curves of the four PP samples were shown in Fig. 9, and each peak of the melting point and the enthalpy of fusion ΔH_m of each sample were listed in Table 4.

It's known that higher melting temperature on the SSA final melting curve was corresponding to higher isotacticity and longer isotactic sequence length in the molecular chains.^{4,7} Fig. 9 and Table 4 showed that there were five melting peaks on the melting curves after SSA fractionation for all PP samples, and especially the higher melt peaks of homo-polypropylene (sample 1 and sample 2) were more different from those of co-polypropylene (sample 3 and sample 4) while the lower melting peaks of polypropylene samples were somewhat similar. Compared with the homo-polypropylene (sample 1 and

Table 4 The SSA results of four PP samples

Samples	$\Delta H_m/(\text{J g}^{-1})$	$T_{m1}/^\circ\text{C}$	$T_{m2}/^\circ\text{C}$	$T_{m3}/^\circ\text{C}$	$T_{m4}/^\circ\text{C}$	$T_{m5}/^\circ\text{C}$
Sample 1	127.1	177.7	170.4	165.8	160.1	154.1
Sample 2	126.2	177.7	170.2	165.7	160.1	154.0
Sample 3	104.5	176.7	171.0	166.3	160.9	154.9
Sample 4	109.7	176.4	172.7	167.8	163.7	158.4

Table 5 The SSA results of four PP samples

Samples	R^{2a}	n_1^b (%)	n_2 (%)	n_3 (%)	n_4 (%)	n_5 (%)
Sample 1	0.981	31.3	36.2	17.1	10.0	5.4
Sample 2	0.977	30.7	36.7	17.5	9.7	5.4
Sample 3	0.985	45.3	23.8	17.8	8.6	4.5
Sample 4	0.996	41.7	32.6	12.4	8.9	4.4

^a The peakfit variance R^2 of the fitted peaks by using Peakfit 4.12 Software. ^b The relative content of peak 1 on the SSA curve of polypropylene calculated using Peakfit 4.12 Software.

sample 2), the molecular chains of the co-polypropylene (sample 3 and sample 4) were more regular while the enthalpy of fusion ΔH_m was lower. In order to quantitatively evaluate the variations of all melting peaks, the relative contents (the integral area percentages on the SSA melting curve) of each peak were calculated using Peakfit 4.12 Software, and the relative contents of all peaks were shown in Table 5. Fig. 10 showed plots of SSA melting curve as well as its fitting curves of sample 1 after treated by Peakfit 4.12 Software. Fig. 11 showed the fitting peaks of SSA melting curves of all PP samples after treatment, and the weight fractions content of the polypropylene fractions calculated by using Peakfit 4.12 Software was shown as Fig. 12.

As can be seen from Table 5 and Fig. 12, the relative contents of peak 1 for the highest isotactic component in the molecular chain of co-polypropylene (sample 3 and sample 4) were about 10–15% higher than that of homo-polypropylene (sample 1 and sample 2), showing that there were more highest isotactic component in the molecular chain of co-polypropylene (sample 3 and sample 4). Table 5 and Fig. 12 also showed that the

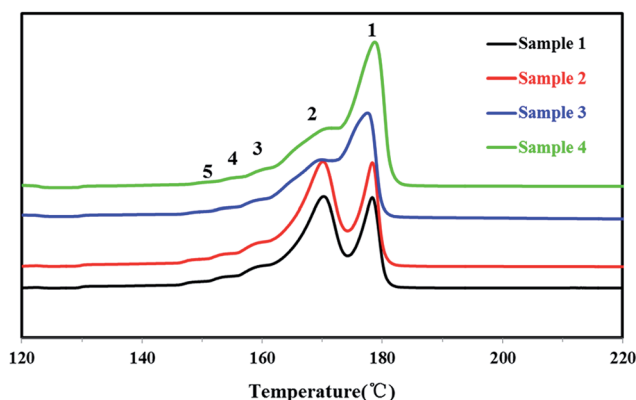


Fig. 9 The SSA melting curves of four PP samples.

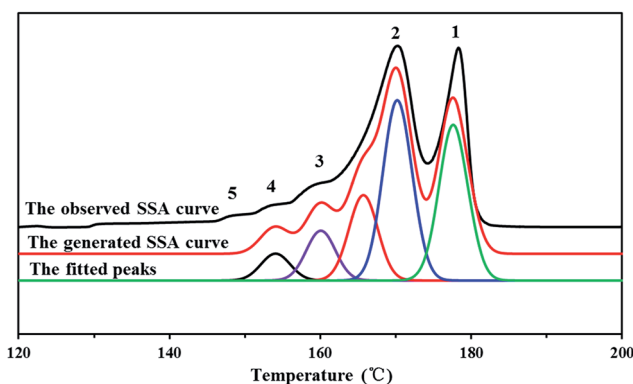


Fig. 10 Plots of SSA melting curve and its fitting curves of PP sample 1 by using Peakfit 4.12 Software where the peakfit variance R^2 of the fitted peaks is 0.981.



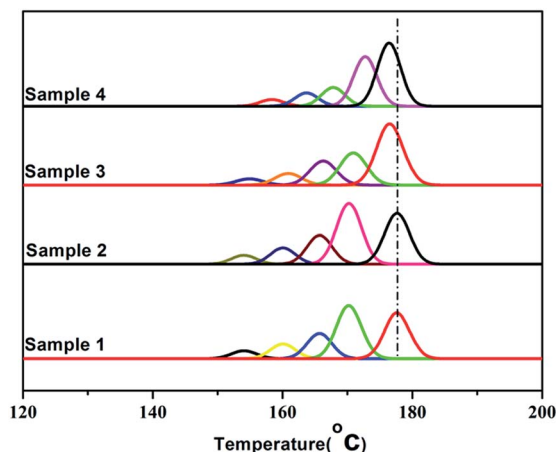


Fig. 11 The fitting curves using Peakfit 4.12 Software of SSA melting curve of PP samples.

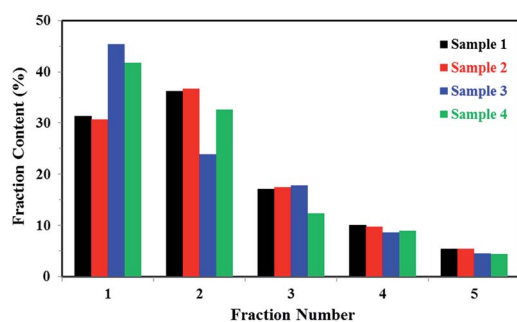


Fig. 12 Weight fractions (%) of the polypropylene fractions calculated by using Peakfit 4.12 Software.

relative contents of (peak 2, peak 3, peak 4, and peak 5) for the higher or lower isotactic component in the molecular chain of co-polypropylene (sample 3 and sample 4) were lower than those of homo-polypropylene (sample 1 and sample 2), showing that the higher or lower isotactic component in the molecular chain of homo-polypropylene (sample 1 and sample 2) were much more than those of co-polypropylene (sample 3 and sample 4). The initial SSA results indicated that the polar copolymer monomer had a great influence on the high isotactic component in co-polypropylene microstructure.

The crystallization sequences with different length of polypropylene could be crystal independently and form the microcrystalline with different sizes when polypropylene was fractionated through thermal techniques such as SSA. Therefore, the crystallization sequence length of polypropylene was related to the lamellar thickness, and the multiplex melting peaks on the SSA melting curves in fact were the melting peaks of the microcrystalline with different sizes formed and annealed at each self-nucleation temperature employed.^{4,7,11,27,46,47} The lamellar thickness could be estimated from the SSA results with the following Thomson–Gibbs equation.⁴⁸

$$T_m = T_m^0 \left(1 - \frac{2\sigma}{\Delta H_0 L_i} \right)$$

where the equilibrium melting temperature $T_m^0 = 460$ K (estimated values between 459–467 K),^{33,49} $\Delta H_0 = 184 \times 10^6 \text{ J m}^{-3}$, the surface energy $\sigma = 0.0496 \text{ J m}^{-2}$ and L_i is the lamellar thickness.⁵⁰

The lamellar thicknesses of all polypropylene samples were calculated by Thomson–Gibbs equation after SSA thermal fractionation, and listed in Table 6.

Table 6 showed the lamellar thicknesses of the highest isotactic component in the molecular chain of homo-polypropylene (sample 1 and sample 2), which were about 3–4 nm longer than those of co-polypropylene (sample 3 and sample 4), showing that the highest isotactic sequence lengths of homo-polypropylene (sample 1 and sample 2) were longer than those of co-polypropylene (sample 3 and sample 4). What is more interesting is that the relative contents of peak 1 for the highest isotactic component in the molecular chain of homo-polypropylene (sample 1 and sample 2) were lower than those of co-polypropylene (sample 3 and sample 4), which showed the microstructure differences in molecular chain of different kind of PP. As the Tables 5 and 6 showed, the lamellar thicknesses of the higher or lower isotactic component in the molecular chain of homo-polypropylene (sample 1 and sample 2) were slightly smaller than those of co-polypropylene (sample 3 and sample 4) while the relative contents of the peaks (peak 2, peak 3, peak 4, and peak 5) for the higher or lower isotactic component in the molecular chain of co-polypropylene (sample 3 and sample 4) of homo-polypropylene (sample 1 and sample 2) were much greater than those of co-polypropylene (sample 3 and sample 4), indicating that the highest isotactic sequence lengths of homo-polypropylene (sample 1 and sample 2) were longer. And the relative contents of high melt peaks ($\geq 160^\circ\text{C}$) of co-polypropylene (sample 3 and sample 4) was very different compared to homo-polypropylene (sample 1 and sample 2), illustrating the polar copolymer monomer had a great influence on the high isotactic component in co-polypropylene microstructure.

Furthermore, the statistical parameters relating to the lamellar thickness, the arithmetic average L_n , weighted average L_w , and the broadness index I , were calculated using the equations as follows:⁵¹

$$L_n = \frac{n_1 L_1 + n_2 L_2 + n_3 L_3 + n_4 L_4 + \dots + n_j L_j}{n_1 + n_2 + n_3 + n_4 + \dots + n_j} = \sum f_i L_i$$

$$L_w = \frac{n_1 L_1^2 + n_2 L_2^2 + n_3 L_3^2 + n_4 L_4^2 + \dots + n_j L_j^2}{n_1 L_1 + n_2 L_2 + n_3 L_3 + n_4 L_4 + \dots + n_j L_j} = \frac{\sum f_i L_i^2}{\sum f_i L_i}$$

Table 6 The lamellar thicknesses of polypropylene samples after SSA thermal fractionation

Samples	L_1/nm	L_2/nm	L_3/nm	L_4/nm	L_5/nm
Sample 1	27.10	15.08	11.78	9.27	7.57
Sample 2	27.10	14.89	11.73	9.27	7.55
Sample 3	24.43	15.65	12.07	9.56	7.76
Sample 4	23.73	17.53	13.02	10.71	8.72



Table 7 Lamellar thickness statistical parameters of PP

Samples	L_n /(nm)	L_w /(nm)	$I = L_w/L_n$
Sample 1	17.29	20.10	1.16
Sample 2	17.14	19.97	1.17
Sample 3	18.11	20.16	1.11
Sample 4	18.56	19.91	1.10

$$I = \frac{L_w}{L_n}$$

where n_i represented as the normalized peak area, and L_i stood for the lamellar thickness for each fraction. The results of all samples were listed in Table 7.

Table 7 showed that the arithmetic average lamellar thicknesses of homo-polypropylene (sample 1 and sample 2) were a little thinner than those of co-polypropylene (sample 3 and sample 4), and the statistical broadness index I of the lamellar thickness distribution of homo-polypropylene samples was wider, suggesting that the isotactic sequence length of homo-polypropylene (sample 1 and sample 2) was not only longer as listed in Table 6, but also the isotactic sequence length distribution of homo-polypropylene samples was broader than that of co-polypropylene samples.

Conclusions

In order to study the microstructure difference between the homo-polypropylene and co-polypropylene, the stereo-defects and its distribution of four homo- and co-polypropylene samples were studied through the correct design SSA, and other characterization ways such as DSC, GPC, and TREF, respectively. Firstly, the preliminary characterization found that the homo-polypropylene or co-polypropylene samples had the similar mechanical and thermal properties, and the SSA results also revealed the internal microstructure of the different kinds of polypropylene was almost exactly the same, representing the similar isotactic sequence length, the fractions weight of the isotactic sequence, and the isotactic sequence length distribution of the same kind of polypropylene. Compared with the homo-polypropylene, the DSC, GPC, and TREF tests indicated there were different properties of the co-polypropylene samples, revealing the differences in molecular chain microstructure. The SSA results after the correction of T_g for each sample displayed there were obvious differences in the (statistical) lamellar thickness and its distribution and the fractions weight of different fractions after SSA treatment, further proving the microstructure difference in the isotactic sequence length, and the isotactic sequence length distribution of homo- and co-polypropylene samples. Besides, the correct design SSA also indicated the polar copolymer monomer had a great influence on the high isotactic component in co-polypropylene microstructure.

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Notes and references

- 1 J. Qiao, M. Guo, L. Wang, D. Liu, X. Zhang, L. Yu, W. Song and Y. Liu, *Polym. Chem.*, 2011, **2**, 1611–1623.
- 2 H. F. Chang, Y. Ma, W. Liang and S. Lai, *RSC Adv.*, 2016, **6**, 26775–26782.
- 3 G. Venugopal, J. Moore, J. Howard and S. Pendalwar, *J. Power Sources*, 1999, **77**, 34–41.
- 4 J. Kang, F. Yang, T. Wu, H. L. Li, D. M. Liu, Y. Cao and M. Xiang, *J. Appl. Polym. Sci.*, 2012, **125**, 3076–3083.
- 5 H. F. Chang, S. T. Ren, T. Zheng, X. F. Dang, Y. Yang, L. Y. Zhang and H. Y. Li, *Acta Polym. Sin.*, 2013, 199–207.
- 6 C. H. Zhang, Y. G. Shangguan, R. F. Chen, Y. Z. Wu, F. Chen, Q. A. Zheng and G. H. Hu, *Polymer*, 2010, **51**, 4969–4977.
- 7 J. Kang, F. Yang, T. Wu, H. L. Li, Y. Cao and M. Xiang, *Eur. Polym. J.*, 2012, **48**, 425–434.
- 8 H. F. Chang, S. T. Ren, X. F. Dang, L. Y. Zhang, H. Y. Li and Y. L. Hu, *J. Appl. Polym. Sci.*, 2013, **129**, 1026–1035.
- 9 J. Kang, Y. Cao, H. Li, J. Li, S. Chen, F. Yang and M. Xiang, *J. Polym. Res.*, 2012, **19**, 1–11.
- 10 C. Zhang, Y. Shangguan, R. Chen and Q. Zheng, *J. Appl. Polym. Sci.*, 2011, **119**, 1560–1566.
- 11 A. J. Müller, Z. H. Hernandez, M. L. Arnal and J. J. Sanchez, *Polym. Bull.*, 1997, **39**, 465–472.
- 12 A. J. Müller, A. T. Lorenzo and M. L. Arnal, *Macromol. Symp.*, 2009, **277**, 207–214.
- 13 A. J. Müller and M. L. Arnal, *Prog. Polym. Sci.*, 2005, **30**, 559–603.
- 14 A. J. Müller, M. L. Arnal, A. L. Spinelli, E. Canizales, C. C. Puig and H. Wang, *Macromol. Chem. Phys.*, 2003, **204**, 1497–1513.
- 15 A. J. Müller, V. Balsamo, M. L. Arnal, T. Jakob, H. Schmalz and V. Abetz, *Macromolecules*, 2002, **35**, 3048–3058.
- 16 M. L. Arnal, E. Canizales and A. J. Müller, *Polym. Eng. Sci.*, 2002, **42**, 2048–2063.
- 17 M. L. Arnal, V. Balsamo, G. Ronca, A. Sanchez, A. J. Müller, E. Canizales and C. U. de Navarro, *J. Therm. Anal. Calorim.*, 2000, **59**, 451–470.
- 18 A. T. Lorenzo, M. L. Arnal, J. J. Sanchez and A. J. Müller, *J. Polym. Sci., Part B: Polym. Phys.*, 2006, **44**, 1738–1750.
- 19 D. Cavallo, A. T. Lorenzo and A. J. Müller, *J. Polym. Sci., Part B: Polym. Phys.*, 2016, **54**, 2200–2209.
- 20 A. T. Lorenzo, M. L. Arnal, A. J. Müller, M. C. Lin and H. L. Chen, *Macromol. Chem. Phys.*, 2011, **212**, 2009–2016.
- 21 R. M. Michell, A. J. Müller, G. Deshayes and P. Dubois, *Eur. Polym. J.*, 2010, **46**, 1334–1344.
- 22 C. J. Perez, N. Villarreal, J. M. Pastor, M. D. Failla, E. M. Valles and J. M. Carella, *Polym. Degrad. Stab.*, 2009, **94**, 1639–1645.
- 23 M. F. J. Pijpers and V. B. F. Mathot, *J. Therm. Anal. Calorim.*, 2008, **93**, 319–327.



- 24 J. Chau and J. Teh, *J. Therm. Anal. Calorim.*, 2005, **81**, 217–223.
- 25 M. Bialek, K. Czaja and B. Sacher-Majewska, *Thermochim. Acta*, 2005, **429**, 149–154.
- 26 V. Virkkunen, P. Laari, P. Pitkänen and F. Sundholm, *Polymer*, 2004, **45**, 3091–3098.
- 27 V. Virkkunen, P. Laari, P. Pitkänen and F. Sundholm, *Polymer*, 2004, **45**, 4623–4631.
- 28 H. F. Chang, H. Y. Li, T. Zheng, Q. Zhou, L. Y. Zhang and Y. L. Hu, *J. Polym. Res.*, 2014, **21**, 1–11.
- 29 J. Kong, X. D. Fan, Y. C. Xie and W. Q. Qiao, *J. Appl. Polym. Sci.*, 2004, **94**, 1710–1718.
- 30 P. Starck, K. Rajanen and B. Lofgren, *Thermochim. Acta*, 2003, **395**, 169–181.
- 31 J. Nicolas, N. Villarreal, I. Gobernado-Mitre, J. C. Merino and J. M. Pastor, *Macromol. Chem. Phys.*, 2003, **204**, 2212–2221.
- 32 Q. Zhou, A. L. Wang, H. Y. Li, Z. Luo, T. Zheng, L. Y. Zhang and Y. L. Hu, *RSC Adv.*, 2016, **6**, 75023–75031.
- 33 E. B. Bond, J. E. Spruiell and J. S. Lin, *J. Polym. Sci., Part B: Polym. Phys.*, 1999, **37**, 3050–3064.
- 34 G. Morini, E. Albizzati, G. Balbontin, I. Mingozzi, M. C. Sacchi, F. Forlini and I. Tritto, *Macromolecules*, 1996, **29**, 5770–5776.
- 35 Y. D. Fan, C. Y. Zhang, Y. H. Xue, X. Q. Zhang, X. L. Ji and S. Q. Bo, *Polymer*, 2011, **52**, 557–563.
- 36 T. Zheng, Q. Zhou, Q. Li, H. Y. Li, L. Y. Zhang and Y. L. Hu, *RSC Adv.*, 2015, **5**, 9328–9336.
- 37 Y. H. Xue, Y. D. Fan, S. Q. Bo and X. L. Ji, *Eur. Polym. J.*, 2011, **47**, 1646–1653.
- 38 I. Suarez, M. J. Caballero and B. Coto, *Eur. Polym. J.*, 2011, **47**, 171–178.
- 39 A. J. Müller, R. M. Michell, R. A. Pérez and A. T. Lorenzo, *Eur. Polym. J.*, 2015, **65**, 132–154.
- 40 B. Fillon, J. C. Wittmann, B. Lotz and A. Thierry, *J. Polym. Sci., Part B: Polym. Phys.*, 1993, **31**, 1383–1393.
- 41 H. F. Chang, Y. Zhang, S. T. Ren, X. F. Dang, L. Y. Zhang, H. Y. Li and Y. L. Hu, *Polym. Chem.*, 2012, **3**, 2909–2919.
- 42 T. F. J. Pijpers, V. B. F. Mathot, B. Goderis, R. L. Scherrenberg and E. W. van der Vegte, *Macromolecules*, 2002, **35**, 3601–3613.
- 43 E. P. Moore, *Polypropylene Handbook: Polymerization, Characterization, Properties, Processing, Applications*, Hanser Publishers, 1996.
- 44 J. Brandrup and E. H. Immergut, *Polymer Handbook*, Wiley, New York, 3rd edn, 1989.
- 45 D. J. Blundell, A. Keller and A. J. Kovacs, *J. Polym. Sci., Part C: Polym. Lett.*, 1966, **4**, 481–486.
- 46 S. J. Song, P. Y. Wu, J. C. Feng, M. X. Ye and Y. L. Yang, *Polymer*, 2009, **50**, 286–295.
- 47 T. Garoff, V. Virkkunen, P. Jaaskelainen and T. Vestberg, *Eur. Polym. J.*, 2003, **39**, 1679–1685.
- 48 U. W. Gedde, *Polymer physics*, Chapman & Hall, London, 1995.
- 49 M. Iijima and G. Strobl, *Macromolecules*, 2000, **33**, 5204–5214.
- 50 A. Wlochowicz and M. Eder, *Polymer*, 1984, **25**, 1268–1270.
- 51 M. Keating, I. H. Lee and C. S. Wong, *Thermochim. Acta*, 1996, **284**, 47–56.

