Synthesis and stereocomplex formation of enantiomeric alternating copolymers with two types of chiral centers, poly(lactic acid-alt-2-hydroxybutanoic acid)s†

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Stereocomplex (SC) formation was reported for the first time for enantiomeric alternating copolymers consisting of repeating units with two types of chiral centers, poly(lactic acid-alt-2-hydroxybutanoic acids [P(LA-alt-2HB)s], L,L'-configured poly(lactic acid-alt-L-2-hydroxybutanoic acid) [PLLAAalt-L-2HB] and d,d'-configured poly(2-hydroxybutanoic acid-alt-2-hydroxybutanoic acid) [PD(LA-alt-2HB)] were amorphous. Blends of PLLAalt-L-2HB and PD(LA-alt-2HB) were crystallizable and showed typical SC-type wide-angle X-ray diffraction profiles similar to those reported for stereocomplexed blends of poly(lactic acid) and poly(o-lactic acid) homopolymers and of poly(L-2-hydroxybutanoic acid) and poly(o-2-hydroxybutanoic acid) homopolymers, and of L,L'-configured poly(lactic acid-co-L-2-hydroxybutanoic acid) [PLLAAco-L-2HB] and D,D'-configured poly(lactic acid-co-O-2-hydroxybutanoic acid) [PD(LA-co-O-2HB)] random copolymers. The melting temperature values and melting enthalpy values at 100% crystallinity for stereocomplexed solvent-evaporated and precipitated PLLAalt-L-2HB/ PD(LA-alt-O-2HB) blends were correspondingly 187.5 and 187.9 °C, and 98.1 and 91.8 J g⁻¹. Enantiomeric polymer blending of PLLAalt-L-2HB and PD(LA-alt-O-2HB) can confer crystallizability by stereocomplexation and the biodegradable materials with a wide variety of physical properties and biodegradability are highly expected to be prepared by synthesis of alternating copolymers of various combinations of two types of chiral α-substituted 2-hydroxalkanoic acid monomers and their SC crystallization.

1. Introduction

Poly(l-lactide) or poly(l-lactic acid) (PLLA), which is produced from renewable plant resources and biodegradable in the environment and the human body, has mechanical properties appropriate for a wide range of applications such as general, biomedical, pharmaceutical, and environmental applications.1-11 To diversify its application, stereocomplex (SC) formation of PLLA with its enantiomer, poly(l-lactide) or poly(o-lactic acid) (PDLA) and α-configured poly(lactide) or poly(lactic acid) (PLA)-like homopolymers and random copolymers has been widely inquired.12-23 Using the method of SC formation between PLLA and PDLA homopolymers and in stereo block copolymers, mechanical properties and resistance to thermal and hydrolytic degradation of PLA-based materials can be achieved.12-21 Furthermore, SC formation of PLA-based block copolymers with water-soluble polymers facilitates the preparation of aqueous biomedical gels.13-25 Various parameters affect SC formation. Such parameters include the types, concentrations and sequence of monomer units,13,14,16,34-39 thermal history,1,18,20,34 external forces,13,18,32-38 the types and concentrations of solvents for casting,16,37-39 the types and concentrations of third polymers and additives,18,40-47

Recently, the symmetry between PLLA and PDLA or their alternating packing was found to be retained in the SC crystal lattice,48,49 whereas PDLA and PLLA chains were reported to be contained in the SC crystal lattice with a wide PLLA fractions from 30 to 70%.50,51 The elastic modulus of the PLLA/PDLA SC crystalline region (20 GPa) in the direction parallel to the c-axis was higher than that of PLLA or PDLA homo-crystalline regions (14 GPa),52 which should have been originated from the stronger interaction between PLLA and PDLA chains than that between

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PLLA chains or PDLA chains, as evidenced by atomic force microscopy.\textsuperscript{24} Also, utilizing hydrogen-bonding, stereo diblock-like PLAs with high stereocomplexationality were prepared from PLLA and PDLA having 2-ureido-[1H]-pyrimidinone terminal,\textsuperscript{44} whereas stereo multiblock PLA was synthesized from DL-lactate at ambient temperature using achiral iron complexes.\textsuperscript{55} The SC formation of lactic acid (LA) and ω-amino acid-based enantiomeric random copoly(esteramides), poly(lactic acid-co-α-alanine) was reported.\textsuperscript{46} Utilizing the stereocomplexed PLA nanofibers, highly transparent self-reinforced PLA composites were prepared.\textsuperscript{57} PLA stereocomplexed materials having different memory shape effects were prepared by network formation or supramolecular formation and incorporation of elastomeric block.\textsuperscript{58,59}

Similar to SC formation between PLLA and PDLA homopolymers, SC formation took place between the enantiomeric PLA-like homopolymers, poly(ω-substituted 2-hydroxyalkanoic acid)s (Fig. 1), such as poly(2-hydroxybutanoic acid) [P(2HB)]\textsuperscript{60} and poly(2-hydroxy-3-methylbutanoic acid) [P(2H3MB)],\textsuperscript{60} and poly(α-mandelic acid).\textsuperscript{40} Hetero SC formation occurred between PLA and P(2HB), P(2HB), and P(2H3MB) with the different chemical structures and opposite configurations. Ternary SC formation took place in the blends of L- and D-configured P(2HB)s and L- or D-configured PLA,\textsuperscript{61} and D-configured P(2HB)s and L- and D-configured P(2H3MB)s.\textsuperscript{62}

Also, SC formation took place in enantiomeric random copolymer blends of poly(l-lactic acid-co-ω-2-hydroxybutanoic acid) [P(LLA-co-ω-2HB)] (56/44) and poly(ω-lactic acid-co-ω-2-hydroxybutanoic acid) [P(DLA-co-ω-2HB)] (52/48)\textsuperscript{63} and of poly(l-lactic acid-co-ω-2-hydroxy-3-methylbutanoic acid) [P(LLA-co-ω-2H3MB)] (47/53) and poly(ω-lactic acid-co-ω-2-hydroxy-3-methylbutanoic acid) [P(DLA-co-ω-2H3MB)] (47/53),\textsuperscript{64} and the staggered random copolymers, L-configured P(LLA-co-L-2HB) (50/50) and D-configured poly(ω-2-hydroxybutanoic acid-co-ω-2-hydroxy-3-methylbutanoic acid) (50/50).\textsuperscript{65} It should be noted that in these cases, all types of monomer units were confirmed to be packed in the SC crystalline lattice by wide-angle X-ray diffractometry (WAXD). This is in marked contrast with normal crystallization of random copolymers, wherein minor monomer units are excluded from and not included in the crystalline regions.

On the other hand, SC formation took place in enantiomeric alternating copolymer blends of poly(l-lactic acid-alt-6-hydroxyhexanoic acid) and poly(ω-lactic acid-alt-6-hydroxyhexanoic acid)\textsuperscript{66} and of poly(l-lactic acid-alt-glycolic acid) [P(LLA-alt-GA)] and poly(ω-lactic acid-alt-glycolic acid) [P(DLA-alt-GA)].\textsuperscript{67} These enantiomeric alternating copolymers which formed SC had the repeating or monomer units of lactic acid-alt-6-hydroxyhexanoic acid or 6-hydroxyhexanoic acid–lactic acid and of lactic acid–glycolic acid or glycolic acid–lactic acid. In these repeating units, only one type of chiral center from lactic acid units existed. On the other hand, Tabata and Abe synthesized alternating copolymers composed of repeating or monomer units of two types of chiral centers, L,D-configured poly(l-lactic acid-alt-ω-3-hydroxybutanoic acid) and L,D-configured poly(l-lactic acid-alt-ω-3-hydroxybutanoic acid) and observed the melting temperature ($T_m$) difference between the two polymers (233 and 83 °C).\textsuperscript{68}

In the present study, we synthesized enantiomeric alternating copolymers of poly(l-lactic acid-alt-ω-2-hydroxybutanoic acid) [P(LLA-alt-ω-2HB)] and poly(ω-lactic acid-alt-ω-2-hydroxybutanoic acid) [P(DLA-alt-ω-2HB)] (Fig. 1) and report SC formation between alternating copolymer blends of P(LLA-alt-ω-2HB) and P(DLA-alt-ω-2HB) at different memory shape effects.
2HB) and P(DLA-alt-D-2HB). These alternating copolymers are composed of repeating unit of lactic acid-2-hydroxybutanoic acid or 2-hydroxybutanoic acid–lactic acid, which contain two types of chiral centers from lactic acid and 2-hydroxybutanoic acid units (Fig. 1). So, this is the first report for SC formation between enantiomeric alternating copolymers consisting of repeating units with two types of chiral centers. To investigate the physical properties, crystalline species, and crystallization behavior of unblended P(LLA-alt-L-2HB) and P(DLA-alt-D-2HB), and their blends, WAXD, differential scanning calorimetry (DSC), and Fourier transform infrared (FTIR) were carried out.

2. Experimental section

Materials and synthesis (Scheme 1),69–72 sample preparation, physical measurements, and observation62,64,67,73–75 were performed according to the literatures. The details of the experiments are stated in the ESI (Section S1).† The weight-average molecular weight (Mw), number average molecular weight (Mn), polydispersity index, and specific optical rotation in chloroform at 25 °C and 589 nm ([α]2559) of P(LLA-alt-L-2HB) and P(DLA-alt-D-2HB) synthesized and used in the present study are 3.00 × 10³, 3.08 × 10³, g mol⁻¹, 1.53 × 10⁴ and 1.54 × 10⁴ g mol⁻¹, 1.96, 2.00, and −122.3, 120.3 deg dm⁻¹ g⁻¹ cm³, respectively.

3. Results and discussion

3.1. Synthesis

In order to corroborate the alternating monomer unit sequences of the synthesized copolymers, 1H and 13C NMR measurements were performed (Fig. 2, S1(a), and S2†). All 1H and 13C NMR spectra for P(LLA-alt-L-2HB) and P(DLA-alt-D-2HB) are shown in ESI as Fig. S1(a) and S2,† respectively, together with those reported for P(2HB), P(LLA-co-2HB) random copolymer, and PLLA73 [Fig. S1(b) and S3†]. The quartet and doublet peaks at 5.2 and 1.6 ppm observed for the 1H NMR spectra of P(LLA-alt-L-2HB) and P(DLA-alt-D-2HB) are attributed to correspondingly methine (b) and methyl (a) protons of lactic acid (LA) units, whereas the quartet, multiplet, and triplet peaks at 5.0, 2.0, and 1.0 ppm are ascribed to correspondingly methine (e), methylene (d), and methyl (c) protons of 2-hydroxybutanoic acid (2HB) units [Fig. S1(a)†], in agreement with the 1H NMR spectrum reported for the random copolymer, P(LLA-co-2HB).75,76 However, methine peaks for LA and 2HB units [(b) and (e)] of P(LLA-alt-L-2HB) are sharp due to their regular alternating monomer sequences, in contrast with broad ones of P(LLA-co-2HB) random copolymer.

The singlet peaks seen at 169.0, 69.0, and 16.8 ppm for 13C NMR spectra of P(LLA-alt-L-2HB) and P(DLA-alt-D-2HB) are attributed to carbonyl, methine, and methyl carbons of LA units, whereas those at 169.9, 73.6, 24.4, and 9.3 ppm are ascribed to carbonyl, methine, methylene, and methyl carbons of 2HB units [Fig. 2(b) and S2†]. In the case of P(LLA-alt-L-2HB) random copolymer, triplet peaks are seen for the methylene carbon of 2HB units (24.2 ppm) and the methine carbon of LA units (16.8 ppm), whereas singlet peaks are observed for P(LLA-alt-L-2HB) and P(DLA-alt-D-2HB) [Fig. 2(b)]. Also, in the cases of P(LLA-alt-L-2HB) and P(DLA-alt-D-2HB), singlet peaks are observed for the carbonyl carbons of 2HB and LA units, whereas in the case of P(LLA-alt-L-2HB) random copolymer, dual peaks are seen for carbonyl carbons of 2HB and LA units [Fig. S2(a) and S3(a)†]. The sharp 1H NMR peaks and singlet 13C NMR peaks of P(LLA-alt-L-2HB) and P(DLA-alt-D-2HB) indicate the fixed circumstances of LA and 2HB units in alternating
sequences but not in the various circumstances in random sequences, confirming the successful synthesis of P(LLA-alt-L-2HB) and P(DLA-alt-D-2HB) alternating copolymers.

3.2. Wide-angle X-ray diffractometry

For estimating the crystalline species and crystallinity \( (X_c) \) values of the samples, WAXD measurements were carried out [Fig. 3(a–c)]. The very weak crystalline peaks seen at 12.8, 19.6, 22.9, and 25.9° for solvent evaporated and precipitated P(DLA-alt-D-2HB) and precipitated P(LLA-alt-L-2HB)/P(DLA-alt-D-2HB) blend are ascribed to those of \( N,N' \)-diisopropylurea (DIU) formed during polymerization and remaining in the samples (Fig. S4†). The unblended P(LLA-alt-L-2HB) and P(DLA-alt-D-2HB) showed broad diffraction profiles, irrespective of sample preparation methods, indicating that the unblended polymer samples were amorphous. This result is in contrast with the results of P(LLA-co-L-2HB) (56/44) and P(DLA-co-D-2HB) (52/48) random copolymers and P(LLA-alt-GA) and P(DLA-alt-GA) alternating copolymers, which were homocrystallizable. The noncrystallizability of P(LLA-alt-L-2HB) and P(DLA-alt-D-2HB) may be due to their lower \( M_w \) values \( (M_w = 3.0 \times 10^3 \text{ and } 3.1 \times 10^3 \text{ g mol}^{-1}, \text{ respectively}) \) compared to those of the P(LLA-co-L-2HB) (56/44) and P(DLA-co-D-2HB) (52/48) random copolymers \( (M_w = 1.4 \times 10^4 \text{ and } 1.6 \times 10^4 \text{ g mol}^{-1}, \text{ respectively}) \) and P(LLA-alt-GA) and P(DLA-alt-GA) alternating copolymers \( (M_w = 4.8 \times 10^3 \text{ and } 5.9 \times 10^3 \text{ g mol}^{-1}, \text{ respectively}) \).

On the other hand, both solvent-evaporated and precipitated P(LLA-alt-L-2HB)/P(DLA-alt-D-2HB) blends showed the crystalline peaks at 11.4° (or 11.2°), 19.8° (or 19.5°), 22.9° (or 22.6°). Here, the data outside and inside parentheses are for the solvent-evaporated and precipitated samples, respectively. The diffraction patterns of P(LLA-alt-L-2HB)/P(DLA-alt-D-2HB) blends are similar to those of PLLA/PDLA homopolymer SC, poly(L-2-hydroxybutanoic acid) [P(L-2HB)]/poly(D-2-hydroxybutanoic acid) [P(D-2HB)] homopolymer SC, and P(LLA-alt-GA) and P(DLA-alt-GA) alternating copolymers SC, but completely different from that of P(LLA-alt-L-2HB)/P(DLA-alt-D-2HB) SC crystallites are shown with broken lines.

**Fig. 2** (a) \(^1\)H NMR spectra and (b) \(^1\)C NMR spectra of methylene carbon of L-2HB units and methyl carbon of LLA units for P(LLA-alt-L-2HB) and P(DLA-alt-D-2HB) alternating copolymers and P(LLA-co-L-2HB) random copolymer in CDCl3. The spectra for P(LLA-co-L-2HB) random copolymer were reproduced from ref. 75 with permission from Elsevier.

**Fig. 3** WAXD profiles of purified, solvent-evaporated, and precipitated (a) P(LLA-alt-L-2HB), and (b) P(DLA-alt-D-2HB), and (c) solvent-evaporated and precipitated P(LLA-alt-L-2HB)/P(DLA-alt-D-2HB) blends. The main diffraction angles of P(LLA-alt-L-2HB)/P(DLA-alt-D-2HB) SC crystallites are shown with broken lines.
crystalline peaks at 17.3°, 18.5°, and 21.3° (ref. 67) [Fig. 4]. The observed 2θ values of P(LLA-alt-L-2HB)/P(DLA-alt-D-2HB) blends are similar to those of P(LLA-co-L-2HB) (56/44)/P(DLA-co-D-2HB) (52/48) random copolymer SC (11.5° (or 11.3°), 20.0° (or 19.6° and 20.1°), 22.9° (or 22.7°), the data outside and inside parentheses are for the solvent-evaporated and melt-crystallized samples, respectively), but lower and higher than those for the melt-crystallized PLLA/PDLA homopolymer SC (11.9°, 20.7°, 24.0°)74 and P(L-2HB)/P(D-2HB) homopolymer SC (10.7°, 18.6° and 19.3°, 21.5°), respectively. The interplanar distance (d) and crystalline diffraction angle (2θ) values of samples were obtained from Fig. 4, except for P(LLA-alt-GA/P(DLA-alt-D-2HB) alternating copolymer SC tabulated in Table S1,† together with those for poly(ε-2-hydroxy-3-methylbutanoic acid) [P(L-2H3MB)]/poly(ε-2-hydroxy-3-methylbutanoic acid) [P(DL-2H3MB)] homopolymer SC. The diffraction peak patterns indicate SC formation between P(LLA-alt-L-2HB) and P(DLA-alt-D-2HB), whereas the 2θ and d values exhibit their SC crystalline lattice sizes are very similar to those of P(LLA-co-L-2HB) (56/44)/P(DLA-co-D-2HB) (52/48) random copolymer SC and between those of PLLA/PDLA homopolymer SC and P(L-2HB)/P(D-2HB) homopolymer SC. Very interestingly, comparison between the WAXD profiles of P(LLA-alt-L-2HB)/P(DLA-alt-D-2HB) SC and P(LLA-alt-GA)/P(DLA-alt-D-2HB) SC [Fig. 4] reveals that their SC crystal lattice types of enantiomeric LA-based alternating copolymer are completely different.

The crystallinity (Xc) values of the samples were obtained from the WAXD profiles in Fig. 3(a–c) and summarized in Table S2.† The Xc values of unblended P(LLA-alt-L-2HB) and P(DLA-alt-D-2HB) (both nil) are much lower than the purified and melt-crystallized unblended P(LLA-co-L-2HB) random copolymer (56/44) (43.7 and 36.5% for samples, respectively)75 and purified and solvent-evaporated unblended P(LLA-alt-GA) and P(DLA-alt-GA) alternating copolymers [8.1% and nil for P(LLA-alt-GA) and 7.2 and 4.8% for P(DLA-alt-GA)]. On the other hand, the solvent-evaporated and precipitated P(LLA-alt-L-2HB)/P(DLA-alt-D-2HB) blends are 66.7 and 60.8%, respectively, which are lower than those reported for the solvent-evaporated and melt-crystallized P(LLA-co-L-2HB) (56/44)/P(DLA-co-D-2HB) (52/48) random copolymer blends (82.5 and 77.7%, respectively), but much higher than those reported for solvent-evaporated and melt-crystallized P(LLA-alt-GA)/P(DLA-alt-GA) alternating copolymer blends (20.8 and 29.2%, respectively). The lower Xc values of unblended P(LLA-alt-L-2HB) and P(DLA-alt-D-2HB) compared to those of unblended P(LLA-co-L-2HB) random copolymer (56/44) and unblended P(LLA-alt-GA) and P(DLA-alt-GA) alternating copolymers and the lower Xc values of P(LLA-alt-L-2HB)/P(DLA-alt-D-2HB) blends compared to those of P(LLA-co-L-2HB) (56/44)/P(DLA-co-D-2HB) (52/48) random copolymer blends can be ascribed to the lower molecular weights of P(LLA-alt-L-2HB) and P(DLA-alt-D-2HB). Interestingly, despite the lower Xc values of unblended P(LLA-alt-L-2HB) and P(DLA-alt-D-2HB) compared to those of unblended P(LLA-alt-GA) and P(DLA-alt-GA) alternating copolymer blends exhibit that dual chiral centers per a repeating unit in P(LLA-alt-L-2HB) and P(DLA-alt-D-2HB) enhanced the crystallizability of the enantiomeric polymers by stereocomplexation compared to one chiral center per a repeating unit in P(LLA-alt-GA) and P(DLA-alt-GA).

3.3. Differential scanning calorimetry

For estimating crystallization and thermal properties of the samples, DSC measurements were conducted (Fig. 5). The glass transition temperature (Tg), Tm, and melting enthalpy (ΔHm) were estimated from the DSC thermograms in Fig. 5 and are summarized in Table S2.† The unblended P(LLA-alt-L-2HB) and P(DLA-alt-D-2HB) show only glass transition peak at around 15.5–40.4 °C without melting peak, regardless of preparation methods, confirming the WAXD result that unblended P(LLA-alt-L-2HB) and P(DLA-alt-D-2HB) were amorphous. On the other hand, the solvent-evaporated and precipitated P(LLA-alt-L-2HB)/P(DLA-alt-D-2HB) blends showed glass transition and melting peaks correspondingly at 33.7, 36.3 °C and 187.5, 187.9 °C. The Tm values for P(LLA-alt-L-2HB)/P(DLA-alt-D-2HB) (Mw = 3.0 × 103 and 3.1 × 103, g mol⁻¹, respectively) blends are higher than those for solvent-evaporated and melt-crystallized (Tc ≈ 70 °C) P(ε-2HB)/P(ε-2HB) (Mw = 1.8 × 103 and 3.3 × 103, g mol⁻¹, respectively) homopolymer blends (Tm = 173.0 and 172.1 °C,
respectively) but lower than those for melt-crystallized \( T_m \) PLLA/PDLA \( (M_w = 4.0 \times 10^3 \text{ g mol}^{-1} \) respectively) homopolymer blends \( (T_m = 197.5 \text{ °C}) \), and solvent-evaporated and melt-crystallized \( (T_c = 160 \text{ °C}) \) P[LLA-co-D-2HB] \( (56/44) \) and P(DLA-co-v-2HB) \( (52/48) \) \( (M_w = 1.4 \times 10^5, 1.6 \times 10^5, \text{ g mol}^{-1} \), respectively) random copolymer blends \( (203.6 \text{ and } 198.4 \text{ °C}) \), and solvent evaporated and melt-crystallized \( (T_c = 100 \text{ °C}) \) P(LLA-alt-GA)/P(DLA-alt-GA) \( (M_w = 4.8 \times 10^3 \text{ and } 5.9 \times 10^3, \text{ g mol}^{-1} \), respectively) blends \( (187.8 \text{ and } 187.6 \text{ °C}) \). The lower \( T_m \) values of P(LLA-alt-L-2HB) and P(DLA-alt-v-2HB) can be attributed to the low molecular weights compared to those of PLLA and PDLA, P(LLA-co-2HB) \( (56/44) \) and P(DLA-co-v-2HB) \( (52/48) \), and P(LLA-alt-GA) and P(DLA-alt-GA).

Using the \( \Delta H_m \) and \( X_c \) of solvent-evaporated \( (65.4 \text{ J g}^{-1} \) and \( 66.7\% \)) and precipitated \( (55.8 \text{ J g}^{-1} \) and \( 60.8\% \)) P(\text{LLA-alt-L-2HB})/P(DLA-co-v-2HB) blends, \( \Delta H_m \) values at \( X_c = 100\% \), i.e., \( \Delta H_m^0 \) values were estimated to be 98.1 and 91.8 J g\(^{-1}\), respectively. Using the \( \Delta H_m \) and \( X_c \) of solvent-evaporated \( (83.7 \text{ J g}^{-1} \) and \( 82.5\% \)) and precipitated \( (74.3 \text{ J g}^{-1} \) and \( 77.7\% \)) P(\text{LLA-co-v-2HB})/P(DLA-co-v-2HB) random copolymer blends, \( \Delta H_m \) values were evaluated to be 101.5 and 95.6 J g\(^{-1}\), respectively. Also, using the \( \Delta H_m \) and \( X_c \) of solvent-evaporated \( (53.4 \text{ J g}^{-1} \) and \( 20.8\% \)) and melt-crystallized \( (67.7 \text{ J g}^{-1} \) and \( 29.2\% \)) P(LLA-alt-GA)/P(DLA-alt-GA) blends, \( \Delta H_m \) values were estimated to be 256.7 and 231.8 J g\(^{-1}\), respectively, which are much higher than \( \Delta H_m^0 \) value of poly(glycolic acid) \( (206 \text{ J g}^{-1}) \), and PLLA/PDLA homopolymer SC \( (142 \text{ (ref. 83) and } 146 \text{ (ref. 77))}) J g\(^{-1}\). Surprisingly, the \( \Delta H_m^0 \) values of P(\text{LLA-alt-L-2HB})/P(DLA-alt-v-2HB) blends are similar to those of P(LLA-co-v-2HB)/P(DLA-co-v-2HB) random copolymer blends, despite the fact that P(\text{LLA-alt-L-2HB}) and P(DLA-alt-v-2HB) have higher sequential regularity compared to that of P(LLA-co-v-2HB) and P(DLA-co-v-2HB). This finding can also be explained by the fact that the molecular weights of P(\text{LLA-alt-L-2HB}) and P(DLA-alt-v-2HB) are lower than those of P(LLA-co-v-2HB) and P(DLA-co-v-2HB) random copolymers. The lower molecular weights or increased density of chain terminals of the former should have increased the defects in the crystalline regions, resulting in lower \( \Delta H_m \) and \( \Delta H_m^0 \) values.

Finally, the present study reveals that even though alternating copolymers with two types of chiral centers per a repeating unit is amorphous, their enantiomeric polymer blending can impose crystallizability by stereocomplexation and it is strongly expected that the biodegradable materials with a wide range of physical properties and biodegradability can be prepared by various combinations of two types of chiral \( \alpha \)-substituted 2-hydroxyalkanoic acid monomers. For this purpose, accumulating the information of physical properties and biodegradability of SCs from enantiomeric alternating \( \alpha \)-substituted hydroxyalkanoic acid-based polymers with various combinations of chiral monomers is required.

4. Conclusions

SC formation was reported for the first time for enantiomeric alternating copolymers with two types of chiral centers per a repeating unit, P(\text{LLA-alt-L-2HB}). \( \alpha \)-configured P(LLA-alt-L-2HB) and \( \nu \)-configured P(DLA-alt-v-2HB) were amorphous polymers. P(\text{LLA-alt-L-2HB})/P(DLA-alt-v-2HB) blends were crystallizable and showed typical SC-type wide-angle X-ray diffraction profiles similar to those reported for PLLA/PDLA homopolymer SC, P(\text{L-2HB})/P(\text{D-2HB}) homopolymer SC, and P(LLA-co-v-2HB)/P(DLA-co-v-2HB) random copolymer SC. The \( T_m \) and \( \Delta H_m \) values for stereocomplexed solvent-evaporated and precipitated P(\text{LLA-alt-L-2HB})/P(DLA-alt-v-2HB) blends were correspondingly 187.5 and 187.9 °C, and 98.1 and 91.8 J g\(^{-1}\). Enantiomeric polymer blending of P(\text{L-alt-L-2HB}) and P(D-alt-v-2HB) can grant crystallizability by stereocomplexation and the biodegradable materials with a wide variety of physical properties and biodegradability are highly expected to be prepared by synthesis of alternating copolymers of various combinations of two types of chiral \( \alpha \)-substituted 2-hydroxyalkanoic acid monomers and their SC crystallization.
Abbreviations

2HB 2-Hydroxybutanoic acid  
ΔH_m Melting enthalpy  
ΔH_m^0 ΔH_m at X_c = 100%  
DIU N,N’-Dioctylpropyleneurea  
DSC Differential scanning calorimetry  
M_n Number average molecular weight  
M_w Weight-average molecular weight  
LA Lactic acid  
P(2HB) Poly(2-hydroxybutanoic acid)  
P(D-2H3MB) Poly(D-2-hydroxy-3-methylbutanoic acid)  
P(PLA-alt-D-2H3MB) Poly(lactic acid-alt-D-2-hydroxy-3-methylbutanoic acid)  
P(LLA-alt-D-2H3MB) Poly(lactic acid-alt-D-2-hydroxy-3-methylbutanoic acid)  
Pr Precipitated  
SE Solvent-evaporated  
[T] Specific optical rotation  
T_g Glass transition temperature  
T_m Melting temperature  
X_c Crystallinity  
WAXD Wide-angle X-ray diffractometry

Conflicts of interest

There are no conflicts to declare.

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