

Cite this: *Polym. Chem.*, 2026, **17**, 923

Synthesis and characterization of biobased (co)polyesters derived from cyclic monomers: camphoric acid and 1,4-cyclohexanedimethanol

Syaiful Ahsan, ^{a,b} Fitrilia Silvianti, ^c Cornelis Post, ^{a,d} Vincent S. D. Voet, ^d Rudy Folkersma, ^d Jeffy Joji, ^e Louis M. Pitet, ^e Subin Damodaran,^f Katja Loos ^a and Dina Maniar ^{*a}

Annual plastic production volumes are more than 400 million tons and are anticipated to continue increasing over the next decade. The majority of plastics originate from fossil resources. Limited raw material reserves and ongoing utilization of plastics contribute to elevated CO₂ emissions, ultimately contributing to climate change. Development of green polymers (*i.e.*, biobased) is one way to reduce our environmental impact. Using renewable resources as raw materials for polymer synthesis reduces the reliance on petroleum and in some cases enables recycling and/or biodegradation. Various aliphatic biobased polyesters have been studied; however, they typically have low glass transition temperatures (T_g) and poor thermomechanical performance, which may limit their applications. In this work, we investigate the synthesis route and structure–property relationships of (co)polyesters from cyclic biobased monomers, camphoric acid and 1,4-cyclohexanedimethanol (CHDM). We observed that increasing the reaction temperature and extending the reaction time led to increased molecular weight and yield of poly(cyclohexanedimethylene camphorate) (PCHC). Conversely, substituting *p*-toluenesulfonic acid (*p*-TSA) with a titanium(IV) isopropoxide (TTIP) catalyst led to reductions in both the molecular weight and yield. Furthermore, (co)polyesters with T_g values ranging from -29 to $+56$ °C were successfully synthesized. DSC and WAXD analyses suggest that the polyesters derived from camphoric acid and the linear diols were amorphous, whereas those based on CHDM were semicrystalline. This work helps address existing knowledge gaps in biobased polymer development by introducing cyclic biobased monomers that expand the current library of renewable materials, thereby broadening opportunities for advanced applications such as coating and packaging materials.

Received 12th December 2025,
Accepted 26th January 2026

DOI: 10.1039/d5py01182e

rsc.li/polymers

Introduction

The predominant monomers utilized in plastic production come from fossil sources, with an increase of global plastic production from 371 million tons (Mt) in 2018 to 414 Mt in 2023, contributing to an increase in CO₂ emissions with its

ongoing use.^{1,2} Currently, one approach to address this issue is the development of green polymers.^{3,4} Green polymer chemistry is characterized by 12 principles, including two primary features: the utilization of renewable resources as raw materials or building blocks for polymer synthesis and the adoption of an environmentally friendly approach for synthesis.⁵ Recently, significant progress has been made in the development of biobased polymers derived from sustainable sources, with increasing interest focused on linear aliphatic polyesters for their biodegradability and on semiaromatic polyesters for their enhanced thermal and mechanical properties. Numerous studies have explored diverse aliphatic biobased monomers and polymerization techniques to optimize their mechanical properties, thermal stability, and degradation behavior, as well as improve the sustainability of the synthetic route.^{6–9} However, these aliphatic polyesters are typically semicrystalline with low glass transition temperatures (T_g) and limited thermomechanical performance, restricting their util-

^aMacromolecular Chemistry & New Polymeric Materials, Zernike Institute for Advanced Materials, University of Groningen, Nijenborgh 3, 9747 AG Groningen, the Netherlands. E-mail: d.maniar@rug.nl

^bPoliteknik STMI Jakarta, the Ministry of Industry of the Republic of Indonesia, Jl. Letjen Suprpto 26, Jakarta, Indonesia

^cLife Science Department, Academy Tech & Design, NHL Stenden University of Applied Sciences, Van Schaikweg 94, 7811 KL Emmen, the Netherlands

^dCircular Plastics, Academy Tech & Design, NHL Stenden University of Applied Sciences, Van Schaikweg 94, 7811 KL Emmen, the Netherlands

^eAdvanced Functional Polymers (AFP) Laboratory, Institute for Materials Research (imo-imomec), Hasselt University, Martelarenlaan 42, 3500 Hasselt, Belgium

^fTosoh Bioscience GmbH, Im Leuschnerpark 4, 64347 Griesheim, Germany



ization for applications demanding high optical clarity, thermal stability, and impact resistance. As a result, designing and synthesizing rigid and amorphous polymers represents a promising strategy for developing biobased materials with enhanced properties.¹⁰

The incorporation of rigid cyclic monomers into polymer backbones is a promising approach to improve the polymer chain stiffness and thermomechanical properties of amorphous polymers. In this context, biobased cyclic monomers have received increasing attention, with camphoric acid from camphor gaining significant interest as a renewable building block for the synthesis of biobased polyesters. Camphor is one of the most widely used commercial aroma chemicals and can be obtained by distilling wood from the camphor laurel tree (*Cinnamomum camphora*).¹¹ It serves as a valuable starting material in the synthesis of a variety of useful chemicals.^{12–14} However, only a limited number of studies have explored the polymerization of camphor-derived monomers.^{15–20} The oxidation of the bicyclic terpene (1*R*)-(+)-camphor results in the production of biobased rigid, five-membered ring diacids known as (1*R*,3*S*)-(+)-camphoric acid.^{15–19} The bioavailability, scalability, and nontoxic nature of camphoric acid make it an appealing platform for sustainable polymer design.²¹ Its rigid structure facilitates the production of polymers with enhanced thermal and mechanical performance, as demonstrated by Ouhichi *et al.*, who developed UV-curable polyester resins from camphoric acid, itaconic acid, and 1,6-hexanediol for coating applications.¹⁶ Guidotti *et al.* synthesized polyester from camphoric acid, butanediol, and another rigid diacid, achieving high barrier properties and mechanical resistance up to 31 MPa, which is suitable for food packaging.^{17,18} Additionally, Hu *et al.* reported a novel biobased epoxy resin from camphoric acid with superior ultraviolet resistance, impact strength, and toughness, offering a sustainable alternative to commercial bisphenol A-based epoxy resins.²⁰ Moreover, Nsengiyumva and Miller demonstrated the degradability of polyesters derived from camphoric acid through water-degradation studies, further emphasizing their potential for sustainable polymer applications.¹⁵

Another cyclic monomer, 1,4-cyclohexanedimethanol (CHDM), has attracted our interest as a valuable cyclic diol for the synthesis of high-performance polyesters. It is traditionally synthesized through a two-step industrial process, wherein dimethyl terephthalate (DMT) is first converted to dimethyl 1,4-cyclohexanedicarboxylate (DMCD), which is subsequently hydrogenated to yield CHDM.²² Alternatively, one-pot direct conversion of DMT to CHDM can be employed.²³ It can also be synthesized from the upcycling of poly(ethylene terephthalate) (PET) waste through tandem reactions utilizing metal catalysts.^{24,25} In an effort to explore possible renewable routes to CHDM, the monomer was synthesized from plant-based acrylate and acetaldehyde, which are alicyclic glycol groups.²⁶ It is commonly used to alter the physical characteristics of PET.²⁷ CHDM was also utilized to increase the melting temperature and glass transition temperature of poly(butylene 2,5-furandicarboxylate) (PBF), a recognized semicrystalline

furanic polyester, which is considered a promising renewable alternative to poly(butylene terephthalate) (PBT) for producing food packaging materials.²⁸ Similar to that of camphoric acid, the rigid structure of CHDM has been reported to promote the production of polyesters, which exhibit high thermomechanical properties and excellent barrier properties.^{29,30} Interestingly, Tsai *et al.* investigated the enzymatic hydrolysis of CHDM-based copolyesters and reported that these polymers undergo surface erosion through random *endo*-type scission, with the rate of degradation being influenced primarily by surface hydrophilicity rather than crystallinity.³¹ Finally, although polymers based on camphoric acid and CHDM have been studied individually, to our knowledge, no investigations have explored polymers derived from their combination. Moreover, a comprehensive understanding of their reaction kinetics, thermomechanical behavior, and biodegradability remains limited. Thus, camphoric acid and CHDM represent promising biobased precursors for synthesizing rigid and amorphous polyesters that combine excellent thermomechanical performance with potential biodegradability.

In this work, we report the synthesis and characterization of biobased (co)polyesters derived from camphoric acid and CHDM. Two aliphatic diols, ethylene glycol (EG) and 1,8-octanediol (OD), as well as cyclic diol CHDM, were used to produce the corresponding polyesters poly(ethylene camphorate) (PEC), poly(octamethylene camphorate) (POC), and poly(cyclohexanedimethylene camphorate) (PCHC), along with two copolyesters: poly(cyclohexanedimethylene-*co*-ethylene camphorate) (P(CHC-*co*-EC)) and poly(cyclohexanedimethylene-*co*-octamethylene camphorate) (P(CHC-*co*-OC)). The variations in these (co)polyesters were explored to understand how the presence of cyclic-aliphatic and cyclic-cyclic polyester structures, as well as their counterpart copolyester polymer chains, influences their properties. The polymerization method was thoroughly investigated, including a kinetic study to examine the distinct catalytic activities and the effects of catalyst variations. To explore potential applications, the correlation between the structure and properties of the obtained polymers was also investigated.

Results and discussion

Synthesis of camphoric acid-based polyesters

Camphoric acid-based polyesters were synthesized *via* a two-step melt polycondensation process—a bulk polymerization method widely favored in industry due to its operational simplicity and efficiency.³² Unlike solvent-based polymerizations, which typically require multiple washing steps and are associated with environmental and economic concerns,³³ the solvent-free approach minimizes such drawbacks. Additionally, reducing or eliminating the solvent content can suppress undesired hydrolysis.³⁴ The synthesis was based on a modified version of the method reported by Nsengiyumva and Miller for PEC,¹⁵ building upon their approach to expand the scope of camphoric acid-based polyesters. Here, the method



was adapted to produce two additional novel polyesters, including one incorporating the cyclic diol CHDM (Table 1). All three camphoric acid-based polyesters were synthesized using a camphoric acid/diol molar ratio of 1.0 : 1.2 (Scheme 1). This ratio was used to compensate for the tendency of diols to evaporate under high-temperature and low-pressure conditions. Polymerization was carried out for 3 or 24 hours at 180 °C in a nitrogen atmosphere, followed by 19 or 7 hours at 2 mmHg when the temperature was increased to 200 or 230 °C. High temperature and vacuum removed water and residual diol, thereby increasing the molecular weight. The products comprised two dark brown sticky polymers (PEC and POC) and one light brown solid polymer (PCHC) (Fig. S12). A summary of the synthesized polyesters is presented in Table 1.

The chemical structures of the obtained polyesters were studied *via* ATR-FTIR and ¹H-NMR spectroscopy. The detailed NMR and IR peak assignments are available in the Experimental section. Fig. 1a shows the ATR-FTIR spectra of the obtained polyesters, confirming the formation of ester linkages through the appearance of an absorption band at 1720–1722 cm⁻¹, which corresponds to the C=O vibration of the ester group. This band represents a shift from the C=O stretching vibration of the carboxylic acid group in camphoric acid (CA), which was originally observed at 1690 cm⁻¹, indicating successful esterification. Additionally, strong absorption bands in the range of 991–1257 cm⁻¹ are observed in all the polyester spectra, corresponding to the asymmetric and symmetric stretching vibrations of C–O bonds. The FTIR spectrum of camphoric acid shows a broad absorption band between 2400–3250 cm⁻¹, attributed to O–H stretching vibrations of hydroxyl groups. This broad band is absent in the spectra of the polyesters, which only show C–H stretching vibrations at 2892–2970 cm⁻¹, further supporting the complete conversion of hydroxyl and carboxylic acid groups into ester functionalities.

The polymeric structure was further confirmed by ¹H-NMR spectroscopy. A representative spectrum of PCHC is shown in Fig. 1b, and those of PEC and POC are provided in the SI (Fig. S1 and S2, respectively). The successful formation of ester linkages is evidenced by the appearance of signals at 4.00 ppm (g – CHDM, CH₂O, *cis*) and 3.91 ppm (g – CHDM, CH₂O, *trans*). These peaks shift downfield from the CH₂OH proton signals of the CHDM monomer, which were originally observed at 3.51 ppm (*cis*-1,4-CHDM) and 3.43 ppm (*trans*-1,4-CHDM) (Fig. S5), consistent with the conversion of the hydroxyl groups into ester functionalities. The residual signals at 3.53 ppm (CH₂OH, *cis*) and 3.46 ppm (CH₂OH, *trans*) are assigned to the end groups (g'), suggesting the presence of hydroxyl functionalities at the polymer chain ends, as previously reported.³⁰

Table 1 lists the abbreviations, polymer structures, molecular weights, dispersities (*D*), and yields of the synthesized polyesters. For polyesters based on linear diol comonomers, PEC and POC, comparable number-average molecular weights (\overline{M}_n) of 14 000 and 14 700 g mol⁻¹, respectively, were obtained. However, POC exhibited a significantly higher weight-average molecular weight (\overline{M}_w), leading to a high dispersity of 3.4, suggesting a broader molecular weight distribution, likely caused by side reactions or non-uniform mixing as the viscosity increased during polymerization. In contrast, the polyester obtained from the polymerization of camphoric acid with the cyclic diol comonomer CHDM (PCHC) had a considerably lower \overline{M}_n value of 4500 g mol⁻¹. The difficulty in removing excess CHDM during condensation, compared to EG or OD, might be a contributing factor. In addition to that, it could also indicate that the bulky structure of cyclic diols (*i.e.*, CHDM) decreases the reaction rate.³⁵ Compared with the work of Nsengiyumva and Miller, who also reported that PEC and related camphoric acid-based polyesters from C₂–C₆ diols with

Table 1 Polymerization conditions and characteristics of the obtained polyesters

Abbreviation	Structure	Reaction conditions ^a	\overline{M}_n^b (g mol ⁻¹)	\overline{M}_w^b (g mol ⁻¹)	<i>D</i>	<i>y</i> ^c (%)
PEC		<i>p</i> -TSA, 180–200 °C, 22 h	14 000	23 200	1.7	43
POC		<i>p</i> -TSA, 180–200 °C, 22 h	14 700	49 500	3.4	63
PCHC		<i>p</i> -TSA, 180–200 °C, 22 h	4 500	8 600	1.9	65
PCHC		<i>p</i> -TSA, 180–230 °C, 31 h	7 800	18 700	2.4	79
PCHC		TTIP, 180–230 °C, 31 h	5 800	13 500	2.3	48

^a Catalyst type, esterification and polycondensation temperature range, and total reaction time. ^b Molecular weights were determined by SEC in chloroform. ^c Isolated yield.





Fig. 2 ^1H -NMR spectra of the mixture from the *p*-TSA-catalyzed polycondensations of camphoric acid and CHDM in bulk, carried out at 180 °C for the first 3 h under nitrogen and increased gradually to 200 °C under dynamic vacuum until 2 mmHg was reached.

duced within 30 minutes of reaction, as evidenced by the emergence of a new signal at 4.00 ppm (*cis*) and 3.91 ppm (*trans*) (**g**, highlighted in light green), corresponding to the methylene group next to the ester bond (COOCH_2). Furthermore, the proton signal attributed to the methylene protons of the hydroxymethyl group (CH_2OH) in the monomer CHDM exhibited a decrease in intensity and a shift from 3.55 ppm (*cis*) and 3.47 ppm (*trans*) to 3.54 ppm (*cis*) and 3.46 ppm (*trans*) (highlighted in light blue), indicating the conversion of the CHDM monomer into oligoesters. This monomer peak disappeared completely after 6 h (3 h vacuum), resulting in the residual signals of the end groups (**g'**). Similarly, protons **b** and **c** in the methyl groups shifted from 0.86 ppm to 0.78 ppm and from 1.25 ppm to 1.21 ppm, respectively. These upfield shifts suggest increased shielding of the protons, likely due to the consumption of electron-withdrawing end groups and their repositioning

within the more electron-rich polymer backbone during the monomer-to-polymer conversion. We also observed an evolution in the ratio between these peaks throughout the reaction time, with the polymer-associated peaks increasing in intensity relative to the monomer peaks, indicating the progressive conversion of monomers into the corresponding polymer. However, when the polymerization was extended from 22 to 26 h, no significant change was observed in the peak intensities, indicating that the reaction was essentially complete. SEC analysis (Table S1 and Fig. S6) corroborated these findings, revealing a gradual increase in molecular weight during the initial 3 h, followed by significant growth after the pressure decreased at 6, 9, and 22 h. Collectively, the ^1H -NMR and SEC results indicate that there is no significant difference between 22, 24, and 26 h, suggesting that a 22 h reaction at 200 °C is sufficient for complete polymerization.





Fig. 3 DSC thermograms of (a) the second heating curve of the PEC, POC, and PCHC and (b) the first heating, cooling, and second heating curves of the PCHC.



Fig. 4 WAXD spectra of (a) poly(ethylene camphorate) (PEC), poly(octamethylene camphorate) (POC) and poly(cyclohexanedimethylene camphorate) (PCHC), and (b) PCHC before and after reheating.

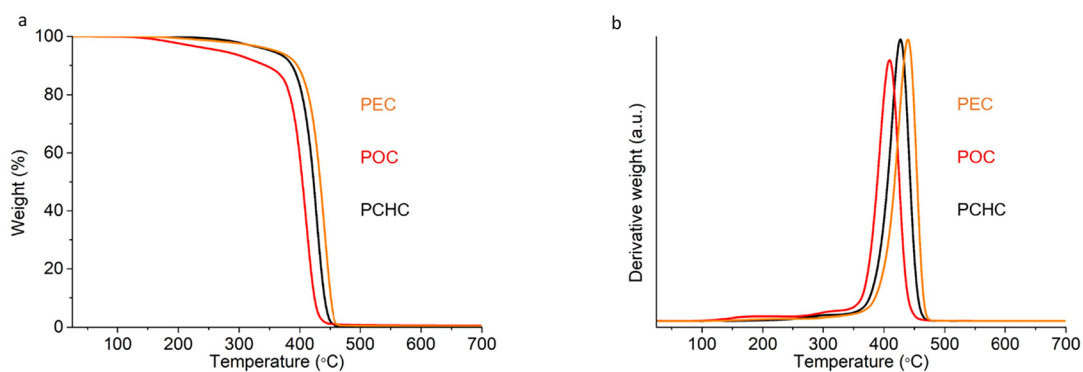


Fig. 5 (a) TGA results of the biobased polyesters: poly(ethylene camphorate) (PEC), poly(octamethylene camphorate) (POC) and poly(cyclohexanedimethylene camphorate) (PCHC). (b) Derivative weight loss curves versus temperature of PEC, POC, and PCHC.

Wettability reflects how strongly a liquid is attracted to a solid surface. The shape of the liquid droplet is determined by the balance between cohesive forces within the liquid and adhesive forces between the liquid and the solid surface.⁵⁴ According to the standard criteria for wettability, systems with a static water contact angle of less than 90° are considered hydrophilic, whereas systems with a contact angle larger than 90° are considered hydrophobic.^{55,56} In this study, static

contact angle measurements were used to evaluate the surface wettability of the synthesized polyester. The contact angles were then determined *via* the drop snake algorithm.⁵⁷ Fig. 6 shows the water contact angles (WCAs) of the CA-derived polyesters. All the measured values are less than 90° , indicating that the polyesters are relatively hydrophilic. PEC and PCHC have similar water contact angle (WCA) values of 82.3° and 87.8° , respectively, whereas POC has a significantly lower WCA





Fig. 6 Drop profile and water contact angle value of (a) PEC, (b) POC, and (c) PCHC.

of 55.5°. This difference may be attributed to the broader molecular weight distribution of POC, as the greater diversity of chain lengths can expose more hydrophilic chain ends, such as hydroxyl (–OH) and carboxylic acid (–COOH) groups. Additionally, surface roughness, which is known to influence water contact angles, should be considered when these results are interpreted.⁵⁷ The incorporation of hydrophilic groups into polymer backbones allows fine-tuning of their wettability, which can, in turn, influence (bio)degradation.⁵⁸ The observed hydrophilicity of PEC, POC, and PCHC suggests their potential for degradation in aqueous environments, which is consistent with preliminary reports showing that polyethylene camphorate (PEC) degrades under both acidic and neutral conditions.¹⁵

Camphoric acid and CHDM-based copolyesters

To further augment the thermomechanical properties of PCHC polyesters, two different series of copolyesters, namely, P(CHC-*co*-EC) and P(CHC-*co*-OC), were synthesized (Table 3). Two different diols, ethylene glycol (EG) or 1,8-octanediol (OD), and CHDM were polymerized with camphoric acid, with varying feed fractions, as listed in Table 4. The polymerization was carried out *via* two-step melt polycondensations for 22 hours at 180–200 °C (Scheme S1). The chemical structures and compositions of both copolyesters were confirmed *via* ¹H-NMR spectroscopy (SI Fig. S3 and S4). The detailed ¹H-NMR peak assignments are available in the Experimental section.

A series of each copolymer was synthesized with three different monomer feed ratios: 25/75, 50/50, and 75/25, corresponding to low, equimolar, and high incorporation of the CHDM monomer, respectively. The resulting copolyesters presented \bar{M}_n values ranging from 5600 to 11 100 g mol^{–1}, with satisfactory yields ranging from 69 to 82% (Table 4). A key

objective of this study was to elucidate the monomer incorporation mechanism during copolymerization, which provides guidance for tuning the thermomechanical properties of the resulting polyesters. We observed that increasing the PCHC molar feed fraction to 50% led to a significant reduction in both the \bar{M}_n and \bar{M}_w values of P(CHC-*co*-EC)s and P(CHC-*co*-OC)s. Further increasing the CHDM content resulted in a subsequent increase in the molecular weight. Although monomer reactivity often influences polycondensations, the close match between the initial feed ratios (F_{CHC} , F_{EC} and F_{OC}) and the resulting copolymer compositions (X_{CHC} , X_{EC} , and X_{OC}) indicates that differences in monomer reactivity do not explain the observed decrease in molecular weight at intermediate CHDM contents. This consistency suggests that all three diol monomers have relatively similar reactivities. At intermediate compositions, the randomness in the copolymer chain sequence often increases, which may reduce the polymerization efficiency and lead to a lower molecular weight and broader dispersity, as evidenced by prior studies.^{30,59,60} This finding is also consistent with our observation that P(CHC-*co*-OC)50 has a broad molecular weight distribution of 3.4. Ultimately, the differences in molar mass are relatively small, considering that small differences in conversion in a step-growth polymerization can lead to rather significant differences in molar mass. In this case, a difference of conversion of less than 2% would account for the discrepancy in molar mass observed by SEC measurements.

Effect of the monomer feed composition on the thermal properties and crystallinity of the copolyesters

The thermal stability of the synthesized copolyesters was evaluated by TGA following the same procedure described above for the camphoric acid-based polyesters. The results are summarized in Table 5 and depicted in Fig. S15. All the copolyesters

Table 3 Polymer structures of copolyesters based on camphoric acid, CHDM and either ethylene glycol or 1,8-octanediol

Abbreviation	Copolyesters	Structure
P(CHC- <i>co</i> -EC)	Poly(cyclohexanedimethylene- <i>co</i> -ethylene camphorate)	
P(CHC- <i>co</i> -OC)	Poly(cyclohexanedimethylene- <i>co</i> -octamethylene camphorate)	



Table 4 Molar fractions, molecular weights, dispersities, and yields of camphoric acid-based copolyesters obtained from various diol feed ratios

Copolymers	Molar fraction (%)				\overline{M}_n^c (g mol ⁻¹)	\overline{M}_w^c (g mol ⁻¹)	<i>D</i>	<i>y</i> ^d (%)
	Feed ^a		Copolyester ^b					
	<i>F</i> _{CHC}	<i>F</i> _{EC}	<i>X</i> _{CHC}	<i>X</i> _{EC}				
P(CHC- <i>co</i> -EC)25	25	75	29	71	8 800	22 800	2.6	77
P(CHC- <i>co</i> -EC)50	50	50	56	44	5 600	14 400	2.6	70
P(CHC- <i>co</i> -EC)75	75	25	78	22	9 500	22 200	2.3	78

Copolymers	<i>F</i> _{CHC}	<i>F</i> _{OC}	<i>X</i> _{CHC}	<i>X</i> _{OC}	\overline{M}_n^c (g mol ⁻¹)	\overline{M}_w^c (g mol ⁻¹)	<i>D</i>	<i>y</i> ^d (%)
	<i>F</i> _{CHC}	<i>F</i> _{OC}	<i>X</i> _{CHC}	<i>X</i> _{OC}				
P(CHC- <i>co</i> -OC)25	25	75	28	72	11 100	28 400	2.6	82
P(CHC- <i>co</i> -OC)50	50	50	49	51	5 700	19 600	3.4	69
P(CHC- <i>co</i> -OC)75	75	25	72	28	8 600	19 800	2.3	81

^a *F*_{CHC}, *F*_{EC}, and *F*_{OC} represent the molar feed ratios of PCHC, PEC, and POC, respectively. ^b *X*_{CHC}, *X*_{EC}, and *X*_{OC} represent the molar fractions of the PCHC, PEC, and POC segments in the obtained copolyesters, respectively, as determined by ¹H-NMR, with calculation details provided in the SI. ^c Molecular weights were determined by SEC in chloroform. ^d Isolated yield.

Table 5 Thermal properties and WAXD data of camphoric acid-based copolyesters

Copolyesters (°C)	DSC ^a		TGA ^b		WAXD ^c		WCA ^d θ (°)
	<i>T</i> _m (°C)	<i>T</i> _g (°C)	<i>T</i> _{d5%} (°C)	<i>T</i> _{dmax}	2θ (°)	χ _c (%)	
P(CHC- <i>co</i> -EC)25	—	17	308	437	—	0	83.3 ± 0.3
P(CHC- <i>co</i> -EC)50	—	29	320	433	—	0	94.1 ± 1.5
P(CHC- <i>co</i> -EC)75	95/106	48	385	426	14.4/16.8	25	101.5 ± 0.4
P(CHC- <i>co</i> -OC)25	—	-18	368	409	—	0	91.2 ± 0.8
P(CHC- <i>co</i> -OC)50	82	2	365	415	14.5/17.1	8	87.4 ± 0.5
P(CHC- <i>co</i> -OC)75	85/109	33	377	420	14.4/16.9	29	105.2 ± 0.3

^a *T*_m (melting temperature) was obtained from the first DSC heating scan; *T*_g (glass transition temperature) was obtained from the second DSC heating scan. ^b *T*_{d5%} (decomposition temperature at 5% weight loss) and *T*_{dmax} (temperature at maximum rate of decomposition) were determined *via* TGA. ^c 2θ (diffraction peak) and χ_c (degree of crystallinity) were determined *via* WAXD. ^d Water contact angle.

exhibited high thermal stability, with *T*_{d5%} values ranging from 308 to 385 °C and *T*_{dmax} values between 409 and 437 °C. Consistent with their corresponding homopolyesters, all *T*_{dmax} values exceeded 400 °C, whereas the char yields remained below 0.62% at 700 °C. The P(CHC-*co*-EC) and P(CHC-*co*-OC) series showed comparable thermal stabilities, with no significant differences between them. This suggests that the incorporation of either C₂ or C₃ aliphatic units has a similar effect on the overall thermal stability of the copolyesters, indicating that the PCHC segments play a dominant role in governing their decomposition behavior. Furthermore, the decomposition temperature (*T*_{d5%} and *T*_{dmax}) did not vary significantly as the PCHC fraction increased from 25 to 75 mol%. This is consistent with previous reports on CHDM-based copolyesters, where the decomposition temperature is largely independent of the monomer feed ratio.²⁹ These results indicate that decomposition is triggered by bond scission, which is not strongly affected by this compositional change.

An increase in the CHDM feed increased the water contact angle from 83.3° to 101.5° for the P(CHC-*co*-EC) series and from 87.4° to 105.2° for the P(CHC-*co*-OC) series. This suggests that the cyclohexanedimethylene-camphorate moiety (the more hydrophobic segment) with lower surface energy, compared with both the ethylene-camphorate and octamethylene-camphorate

moieties, segregated to the surface during spin-coating.^{61,62} Moreover, higher crystallinity increases the molecular mobility of the low surface energy component.^{61,63} However, in the latter series, P(CHC-*co*-OC)50 decreased (87.4°), which can be explained by its broad dispersity (3.4), which is equal to that of POC. The greater variety in the length of the chains would expose their hydrophilic chain ends. This results in the hydrophobicity of polymers that contain a minimum of 50% CHDM (except for P(CHC-*co*-OC)50), perhaps slightly reducing their potential susceptibility to hydrolytic degradation. This remains to be seen in a future extension of this work.

The thermophysical behavior of the copolyester series was analyzed *via* DSC, and the *T*_g and *T*_m values are summarized in Table 5. For the P(CHC-*co*-EC) series, the *T*_g ranged from +17 to +48 °C, whereas for the P(CHC-*co*-OC) series, they ranged from -18 to +33 °C. In both series, the *T*_g values increased with increasing CHDM content, indicating increased chain rigidity and reduced mobility due to the presence of bulky, rigid PCHC segments. These results are consistent with prior research on CHDM-containing copolyesters.^{31,64,65} For example, Diao *et al.* reported that by utilizing CHDM, 1,4-butanediol and furandicarboxylic acid, the *T*_g of polyesters increased from +40 to +87 °C with increasing CHDM feed from 0 to 100%.



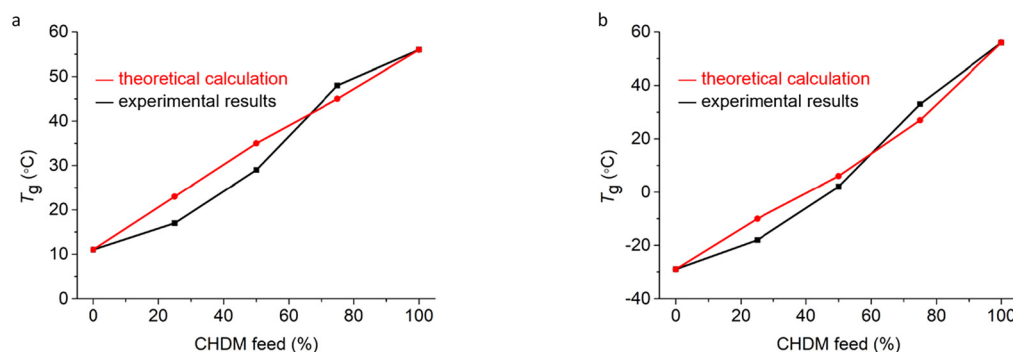


Fig. 7 Comparison of theoretical and experimental glass transition temperatures for (a) P(CHC-co-EC) and (b) P(CHC-co-OC) copolyesters.

All the copolyesters had only one T_g value, which suggests a uniform amorphous phase. The experimentally obtained T_g values were compared with theoretically derived T_g values on the basis of the T_g values of the homopolyesters PEC, POC, and PCHC (Table 2). The theoretical values of the glass transition temperatures of the copolyesters were calculated *via* the Fox equation, as described in the Experimental section.^{31,65–70} A comparison between the theoretical and experimental values is shown in Fig. 7. We observed that there was no significant difference between the theoretical values and the experimental data regarding the T_g values of the copolyesters, suggesting that the copolyesters are random copolymers with good thermodynamic compatibility.^{69,70}

T_m is a first-order transition associated with the crystalline domains of the polymer and was observed only in P(CHC-co-EC)75, P(CHC-co-OC)50, and P(CHC-co-OC)75 (Table 5 and Fig. S14). WAXD results further confirmed the semicrystallinity of these copolyesters, showing sharp peaks at $2\theta \approx 14^\circ$ and 17° , corresponding to their crystalline domains (Fig. S10). In the P(CHC-co-EC)75 sample, two melting endotherms were detected at 95°C and 106°C , indicating that semicrystallinity develops only when the PCHC segment content reaches 75 mol%. In contrast, in the P(CHC-co-OC) series, P(CHC-co-OC)50 already exhibited melting transitions at 82°C , and P(CHC-co-OC)75 did so at 85°C and 109°C , indicating that a lower PCHC fraction (50 mol%) is sufficient to induce crystallinity in this series. This difference suggests that longer octamethylene segments facilitate more effective packing of CHDM-rich regions, enabling crystallization at lower PCHC contents than shorter ethylene segments do. Both crystallinity and surface hydrophilicity are significant factors influencing polymer biodegradation and should be considered, among other factors.⁷¹

Conclusions

In summary, a series of novel (co)polyesters were synthesized in bulk from the rigid, cyclic, biobased monomers camphoric acid and CHDM. Copolymerization of camphoric acid with aliphatic diols and/or CHDM yielded copolyesters with molecular

weights up to $14\,700\text{ g mol}^{-1}$. The polycondensations of camphoric acid and CHDM were investigated under various conditions, including different reaction times, temperatures, and the use of two catalysts. The results showed that using *p*-TSA with a 22 h reaction at 200°C was sufficient to achieve complete polycondensations of camphoric acid and CHDM. The rigid structure of CHDM resulted in an increase in the T_g from 29°C for POC to $+56^\circ\text{C}$ for PCHC, demonstrating that its incorporation as a cyclic comonomer enhances the thermal properties of the polyesters. This trend was further corroborated in the copolyester series: as the PCHC content increased, the T_g values shifted consistently higher, reflecting the restricted chain mobility imparted by the cyclic comonomer. These results demonstrate that T_g can be effectively tuned by adjusting the CHDM incorporation ratio. Moreover, the observed T_g values are in good agreement with theoretical predictions based on the Fox equation.

DSC and WAXD analyses revealed that the incorporation of CHDM promoted semicrystalline behavior in the camphoric acid-based polyesters. For example, PCHC exhibited melting peaks at 99°C and 162°C and a degree of crystallinity of 46%. In the case of the copolyesters, the effect of CHDM content on crystallization is also evident: P(CHC-co-EC)75 shows two melting endotherms at 95°C and 106°C , whereas P(CHC-co-OC)50 and P(CHC-co-OC)75 display melting peaks at $82\text{--}85^\circ\text{C}$ and 109°C , reflecting increasing crystallinity with increasing CHDM content. These results demonstrate that higher CHDM incorporation enhances chain regularity, promoting semicrystallinity and improving thermal stability, whereas camphoric acid alone predominantly forms amorphous polymers.

All the homopolyesters and the two copolyesters are hydrophilic, with water contact angles ranging from 55.5 to 87.8° , indicating favorable interactions with aqueous environments and potential susceptibility to hydrolytic degradation. All the obtained polyesters exhibited high thermal stability, with T_{dmax} values exceeding 400°C and minimal char formation ($<0.6\%$ at 700°C), demonstrating robust polymer backbones capable of withstanding elevated temperatures. These results suggest that the structural durability and environmental responsiveness of polyesters derived from camphoric acid and CHDM are balanced, although further investigations into their



degradability and their long-term performance are still necessary. Finally, these findings demonstrate that biobased rigid monomers such as camphoric acid and CHDM can be effectively utilized to synthesize (co)polyesters with tunable thermal and physicochemical properties, making them promising candidates for applications such as coatings, packaging, and other sustainable materials.

Experimental section

Materials

(1*R*,3*S*)-(+)-Camphoric acid (CA, 99%), ethylene glycol (EG, EMPLURA®), 1,8-octanediol (OD, 98%), 1,4-cyclohexane dimethanol (CHDM, mixture of *cis* and *trans*, 99%), *p*-toluene-sulfonic acid (*p*-TSA, monohydrate, ≥98%), titanium(IV) isopropoxide (TTIP, 97%), chloroform (CHCl₃, amylene stabilized, HPLC grade, >99.8%), and deuterated chloroform (CDCl₃, 99.8 atom% D) were purchased from Sigma-Aldrich. Chloroform (CHCl₃, ChromAR HPLC, ethanol stabilized) and methanol (MeOH, AR®, anhydrous) were purchased from Macron Fine Chemicals. All chemicals were used as received.

General synthetic procedure for the synthesis of camphoric acid-derived biobased (co)polyesters

The synthesis route for camphoric acid and CHDM-based (co)polyesters was modified on the basis of the work of Nsengiyumva and Miller.¹⁵ A two-stage melt polycondensation method was applied. The typical reaction setup is carried out in a 50 ml round-bottom flask equipped with a magnetic stirring egg and connected to a Schlenk line to allow switching between an inert argon flow and high vacuum (2 mmHg). The molar ratio of CA to diol (EG, OD, and CHDM) in the reaction mixture was maintained at 1.0/1.2. The total mass of the monomers was 2.0 g, with an additional 50.0 mg of *p*-TSA used as the catalyst. As an example, the synthesis of poly(ethylene camphorate) (PEC) is as follows: 1.458 g CA (7.282 mmol), 0.542 g CHDM (8.732 mmol), and 50.0 mg *p*-TSA were added to the flask. During the initial 3 h period, the temperature was adjusted to 180 °C while nitrogen was used to create an inert atmosphere. In the second stage of the experiment, the temperature was increased to 190 °C, and a dynamic vacuum was employed until it reached a pressure of 2 mmHg for 19 hours while the temperature was increased to 200 °C.

The synthesis of poly(cyclohexanedimethylene camphorate) (PCHC) required two additional extended procedures, which were adjusted from the previous procedure. In one case, a nitrogen atmosphere was applied for 24 h at 180 °C, followed by 200–230 °C for 7 h in dynamic vacuum until 2 mmHg was reached. In another case, when a similar procedure was used, the catalyst was changed to TTIP.

The polyester products were dissolved in chloroform and filtered, and then the polyesters were precipitated in cold methanol. The precipitation products were centrifuged and decanted, and the obtained polyesters were dried in a vacuum

oven at 40 °C for 72 hours and characterized by ¹H-NMR, FTIR, SEC, DSC, TGA, WAXD, and WCA.

Camphoric acid and camphoric acid-derived biobased (co)polyesters

ATR-FTIR (ν , cm⁻¹): camphoric acid. 2400–3250 (O–H stretching vibration); 2892–2970 (C–H stretching vibrations); 1690 (C=O stretching vibrations); 1411–1458 (C–H bending vibration); 1120–1280 (C–O stretching vibrations); 933 (O–H bending vibrations). Camphoric acid-based (co)polyesters. 2854–2966 (C–H stretching vibrations); 1720–1722 (C=O stretching vibrations); 1353–1460 (C–H bending and stretching vibrations); 991–1257 (C–O stretching vibrations); 754 (C–H bending (rocking) vibrations in POC which possesses more than four CH₂ groups in an aliphatic chain).

Poly(ethylene camphorate) [PEC]. The ¹H-NMR (600 MHz, CDCl₃), δ (ppm): 4.28 (m, 4H, ester–CH₂–CH₂–ester), 3.82 (t, 2H, CH₂–OH end group), 2.78 (t, 1H, CH cyclopentane), 2.53 (m, 1H, CH₂ cyclopentane), 2.15 (m, 1H, CH₂ cyclopentane), 1.81 (m, 1H, CH₂ cyclopentane), 1.50 (m, 1H, CH₂ cyclopentane), 1.23 (s, 3H, CH₃), 1.19 (s, 3H, CH₃), 0.77 (s, 3H, CH₃).

Poly(octamethylene camphorate) [POC]. ¹H-NMR (600 MHz, CDCl₃), δ (ppm): 4.06 (m, 4H, ester–CH₂–CH₂–(CH₂)₄–CH₂–CH₂–ester), 3.38 (t, 2H, (CH₂)₇–CH₂–OH end group), 2.78 (t, 1H, CH cyclopentane), 2.56 (m, 1H, CH₂ cyclopentane), 2.16 (m, 1H, CH₂ cyclopentane), 1.80 (m, 1H, CH₂ cyclopentane), 1.68–1.52 (m, 4H, ester–CH₂–CH₂–(CH₂)₄–CH₂–CH₂–ester), 1.49 (m, 1H, CH₂ cyclopentane), 1.32 (m, 8H, ester–CH₂–CH₂–(CH₂)₄–CH₂–CH₂–ester), 1.25 (s, 3H, CH₃), 1.20 (s, 3H, CH₃), 0.77 (s, 3H, CH₃).

Poly(cyclohexanedimethylene camphorate) [PCHC]. ¹H-NMR (600 MHz, CDCl₃), δ (ppm): 4.00 (m, 2H, ester–CH₂–cyclohexane, *cis*), 3.91 (m, 2H, ester–CH₂–cyclohexane, *trans*), 3.53 (d, 2H, cyclohexane–CH₂–OH end group, *cis*), 3.46 (d, 2H, cyclohexane–CH₂–OH end group, *trans*), 2.79 (t, 1H, CH cyclopentane), 2.57 (m, 1H, CH₂ cyclopentane), 2.17 (m, 1H, CH₂ cyclopentane), 1.91–1.76 (m, 1H, CH₂ cyclopentane, 2H, CH cyclohexane, 2H, CH₂ cyclohexane), 1.65–1.40 (m, 1H, CH₂ cyclopentane, 4H, CH₂ cyclohexane), 1.26 (s, 3H, CH₃), 1.21 (s, 3H, CH₃), 1.04 (m, 2H, CH₂ cyclohexane), 0.77 (s, 3H, CH₃).

Poly(cyclohexanedimethylene-co-ethylene camphorate) [P(CHC-co-EC)]. ¹H-NMR (600 MHz, CDCl₃), δ (ppm): 4.29 (m, 4H, ester–CH₂–CH₂–ester), 3.99 (m, 2H, ester–CH₂–cyclohexane, *cis*), 3.89 (m, 2H, ester–CH₂–cyclohexane, *trans*), 3.61 (t, 2H, CH₂–OH end group), 3.33 (d, 2H, CH₂–OH end group, *cis*), 3.25 (d, 2H, CH₂–OH end group, *trans*), 2.79 (m, 1H, CH cyclopentane), 2.56 (m, 1H, CH₂ cyclopentane), 2.17 (m, 1H, CH₂ cyclopentane), 1.90–1.74 (m, 1H, CH₂ cyclopentane, 2H, CH cyclohexane, 2H, CH₂ cyclohexane), 1.66–1.33 (m, 1H, CH₂ cyclopentane, 4H, CH₂ cyclohexane), 1.25 (s, 3H, CH₃), 1.20 (s, 3H, CH₃), 1.03 (m, 2H, CH₂ cyclohexane), 0.78 (s, 3H, CH₃).

Poly(cyclohexanedimethylene-co-octamethylene camphorate) [P(CHC-co-OC)]. ¹H-NMR (600 MHz, CDCl₃), δ (ppm): 4.06 (m, 4H, ester–CH₂–CH₂–(CH₂)₄–CH₂–CH₂–ester), 4.00 (m, 2H, ester–CH₂–cyclohexane, *cis*), 3.92 (m, 2H, ester–CH₂–cyclohexane, *trans*), 3.37 (m, 2H, (CH₂)₇–CH₂–OH end group), 3.28



editing. K. Loos: supervision, resources, project administration, funding acquisition, writing – review & editing. D. Maniar: conceptualization, methodology, formal analysis, supervision, resources, project administration, funding acquisition, writing – review & editing.

Conflicts of interest

There are no conflicts to declare.

Data availability

The data supporting this article have been included as part of the supplementary information (SI). Supplementary information: spectra of PEC, POC, PCHC, P(CHC-co-EC), P(CHC-co-OC), and CHDM in CDCl₃ (Fig. S1–S5). Molecular weights and SEC chromatograms of the mixture from the *p*-TSA-catalyzed polycondensations of camphoric acid and CHDM in bulk (Table S1 and Fig. S6). SEC chromatograms of PEC, POC, and PCHC (Fig. S7). SEC chromatograms of PCHC from variation of polymerization conditions (Fig. S8). SEC chromatograms of P(CHC-co-EC) and P(CHC-co-OC) (Fig. S9). WAXD patterns of P(CHC-co-EC) and P(CHC-co-OC) (Fig. S10). Multi-peak fitting of WAXD spectra of PCHC, P(CHC-co-EC)75, P(CHC-co-OC)50, and P(CHC-co-OC)75 (Fig. S11). Product appearance of PEC, POC, and PCHC (Fig. S12). Product appearances of P(CHC-co-EC)25, P(CHC-co-EC)50, P(CHC-co-EC)75, P(CHC-co-OC)25, P(CHC-co-OC)50, and P(CHC-co-OC)75 (Fig. S13). DSC thermograms of P(CHC-co-EC)25, P(CHC-co-EC)50, P(CHC-co-EC)75, P(CHC-co-OC)25, P(CHC-co-OC)50, and P(CHC-co-OC)75 (Fig. S14). TGA results of P(CHC-co-EC)s and P(CHC-co-OC)s (Fig. S15). Drop profile and water contact angle value of P(CHC-co-EC)25, P(CHC-co-EC)50, P(CHC-co-EC)75, P(CHC-co-OC)25, P(CHC-co-OC)50, and P(CHC-co-OC)75 (Fig. S16). See DOI: <https://doi.org/10.1039/d5py01182e>.

Acknowledgements

Jur van Dijken is gratefully acknowledged for his assistance with the TGA measurements. The first author (S. Ahsan) expresses gratitude for the financial assistance provided by the Indonesia Endowment Funds for Education of the Ministry of Finance (LPDP Kemenkeu) of the Republic of Indonesia.

References

- M. Zhu and Q. Huang, *Front. Environ. Sci.*, 2025, **13**, 1603040.
- B. M. Stadler, A. Brandt, A. Kux, H. Beck and J. G. de Vries, *ChemSusChem*, 2020, **13**, 556–563.
- R. Mülhaupt, *Macromol. Chem. Phys.*, 2013, **214**, 159–174.
- S. Nandy, E. Fortunato and R. Martins, *Prog. Nat. Sci.: Mater. Int.*, 2022, **32**, 1–9.
- S. Kobayashi, *Struct. Chem.*, 2017, **28**, 461–474.
- Y. Jiang, A. J. J. Woortman, G. O. R. Alberda van Ekenstein and K. Loos, *Biomolecules*, 2013, **3**, 461–480.
- Y. Jiang, G. O. R. A. Van Ekenstein, A. J. J. Woortman and K. Loos, *Macromol. Chem. Phys.*, 2014, **215**, 2185–2197.
- D. Maniar, C. Fodor, I. K. Adi, A. J. J. Woortman, J. van Dijken and K. Loos, *Polym. Int.*, 2021, **70**, 555–563.
- P. Shan, X. Lian, W. Lu, X. Yin, Y. Lu, M. Zhang, X. Wen, G. Xin, Z. Li and Z. Li, *Biomacromolecules*, 2023, **24**, 2563–2574.
- N. G. Valsange, M. N. Garcia Gonzalez, N. Warlin, S. V. Mankar, N. Rehnberg, S. Lundmark, B. Zhang and P. Jannasch, *Green Chem.*, 2021, **23**, 5706–5723.
- S. Guo, Z. Geng, W. Zhang, J. Liang, C. Wang, Z. Deng and S. Du, *Int. J. Mol. Sci.*, 2016, **17**, 1836.
- Y. Zhang, P. Zhao, S. Sun, Q. Wu, E. Shi and J. Xiao, *Commun. Chem.*, 2023, **6**, 133.
- Z. Rafiński and A. Kozakiewicz, *J. Org. Chem.*, 2015, **80**, 7468–7476.
- S. T. Duong and M. Fujiki, *Polym. Chem.*, 2017, **8**, 4673–4679.
- O. Nsengiyumva and S. A. Miller, *Green Chem.*, 2019, **21**, 973–978.
- R. Ouhichi, A. Bougarech, M. Kluge, S. Pérocheau Arnaud, S. Abid, M. Abid and T. Robert, *Eur. Polym. J.*, 2021, **151**, 110423.
- G. Guidotti, M. Soccio, M. Gazzano, V. Siracusa and N. Lotti, *Molecules*, 2023, **28**, 4056.
- G. Guidotti, G. Burzotta, M. Soccio, M. Gazzano, V. Siracusa, A. Munari and N. Lotti, *Polymers*, 2021, **13**, 2707.
- C. Pang, X. Jiang, Y. Yu, L. Chen, J. Ma and H. Gao, *ACS Macro Lett.*, 2019, **8**, 1442–1448.
- J. Hu, C. Wang, J. Dai, N. Teng, S. Wang, L. Zhang, Y. Jiang and X. Liu, *Polym. Adv. Technol.*, 2021, **32**, 3701–3713.
- C. E. Carraher, M. R. Roner, A. G. Campbell, A. Moric-Johnson, L. Miller, P. Slawek, F. Mosca, J. D. Einkauf, J. E. Haky and R. Crichton, *J. Inorg. Organomet. Polym. Mater.*, 2018, **28**, 481–491.
- S. R. Turner, *J. Polym. Sci., Part A: Polym. Chem.*, 2004, **42**, 5847–5852.
- X. Xiao, H. Xin, Y. Qi, C. Zhao, P. Wu and X. Li, *Appl. Catal., A*, 2022, **632**, 118510.
- W. Ren, C. Gang, C. Zhao, P. Wu and X. Li, *Appl. Catal., A*, 2025, **698**, 120233.
- S. Mo, J. Kou, J. Zeng, K. Song, Y. Zhang, S. He, Y. Hu, Y. Guo, X. Liu, X. Chen and Y. Wang, *Chem. Eng. J.*, 2024, **500**, 157249.
- L. Yuan, Y. Hu, Z. Zhao, G. Li, A. Wang, Y. Cong, F. Wang, T. Zhang and N. Li, *Angew. Chem., Int. Ed.*, 2022, **61**, e202113471.
- H. J. Bang, H. Y. Kim, F. L. Jin and S. J. Park, *J. Ind. Eng. Chem.*, 2011, **17**, 805–810.
- L. Diao, K. Su, Z. Li and C. Ding, *RSC Adv.*, 2016, **6**, 27632–27639.
- J. Wang, X. Liu, Y. Zhang, F. Liu and J. Zhu, *Polymer (Guildf.)*, 2016, **103**, 1–8.



- 30 L. Urpí, A. Alla and A. Martínez de Ilarduya, *Polymer (Guildf.)*, 2023, **266**, 125624.
- 31 Y. Tsai, L. C. Jheng and C. Y. Hung, *Polym. Degrad. Stab.*, 2010, **95**, 72–78.
- 32 C. Post, J. van der Vlist, J. A. Jongstra, R. Folkersma, V. S. D. Voet and K. Loos, *Eur. Polym. J.*, 2025, **222**, 113594.
- 33 I. K. Park, H. Sun, S. H. Kim, Y. Kim, G. E. Kim, Y. Lee, T. Kim, H. R. Choi, J. Suhr and J. Do Nam, *Sci. Rep.*, 2019, **9**, 7033.
- 34 A. K. Chaudhary, J. Lopez, E. J. Beckman and A. J. Russell, *Biotechnol. Prog.*, 1997, **13**, 318–325.
- 35 W. J. Yoon, S. Y. Hwang, J. M. Koo, Y. J. Lee, S. U. Lee and S. S. Im, *Macromolecules*, 2013, **46**, 7219–7231.
- 36 Z. Terzopoulou, E. Karakatsianopoulou, N. Kasmi, M. Majdoub, G. Z. Papageorgiou and D. N. Bikiaris, *J. Anal. Appl. Pyrolysis*, 2017, **126**, 357–370.
- 37 Z. Terzopoulou, E. Karakatsianopoulou, N. Kasmi, V. Tsanaktis, N. Nikolaidis, M. Kostoglou, G. Z. Papageorgiou, D. A. Lambropoulou and D. N. Bikiaris, *Polym. Chem.*, 2017, **8**, 6895–6908.
- 38 J. Zhu, J. Cai, W. Xie, P. H. Chen, M. Gazzano, M. Scandola and R. A. Gross, *Macromolecules*, 2013, **46**, 796–804.
- 39 A. G. Amador, A. Watts, A. E. Neitzel and M. A. Hillmyer, *Macromolecules*, 2019, **52**, 2371–2383.
- 40 M. S. Mohammadnia, P. Salaryan, Z. K. Azimi and F. T. Seyidov, *Int. J. Chem. Biochem. Sci.*, 2012, **2**, 36–41.
- 41 C. Q. Huang, S. Y. Luo, S. Y. Xu, J. B. Zhao, S. L. Jiang and W. T. Yang, *J. Appl. Polym. Sci.*, 2010, **115**, 1555–1565.
- 42 T. Zhang, B. A. Howell, P. K. Martin, S. J. Martin and P. B. Smith, *Green Polym. Chem.*, 2013, **1144**, 281–290.
- 43 V. S. R. Ganga, S. H. R. Abdi, R. I. Kureshy, N. U. H. Khan and H. C. Bajaj, *J. Mol. Catal. A:Chem.*, 2016, **420**, 264–271.
- 44 K. S. Naidu, A. S. Kate, V. Kshirsagar, R. Ganeshan, T. Gunale, B. Zhou, S. Anapat, Y. Sulub, A. Kumar and N. Rao, *Polym. Test.*, 2020, **86**, 106446.
- 45 J. Wang, X. Liu, Z. Jia, L. Sun, Y. Zhang and J. Zhu, *Polymer (Guildf.)*, 2018, **137**, 173–185.
- 46 S. Curia, A. Biundo, I. Fischer, V. Braunschmid, G. M. Gübitz and J. F. Stanzione, *ChemSusChem*, 2018, **11**, 2529–2539.
- 47 P. Pan and Y. Inoue, *Prog. Polym. Sci.*, 2009, **34**, 605–640.
- 48 M. C. Righetti, P. Marchese, M. Vannini, A. Celli, C. Lorenzetti, D. Cavallo, C. Ocando, A. J. Müller and R. Androsch, *Biomacromolecules*, 2020, **21**, 2622–2634.
- 49 Y. T. Shieh and G. L. Liu, *J. Polym. Sci., Part B:Polym. Phys.*, 2007, **45**, 466–474.
- 50 G. Stoclet, R. Seguela and J. M. Lefebvre, *Polymer (Guildf.)*, 2011, **52**, 1417–1425.
- 51 E. Núñez, C. Ferrando, E. Malmström, H. Claesson, P. E. Werner and U. W. Gedde, *Polymer (Guildf.)*, 2004, **45**, 5251–5263.
- 52 D. C. Bassett, R. H. Olley, I. A. M. Ai and R. J. J. Thomson, *Polymer (Guildf.)*, 1988, **29**, 1745–1754.
- 53 G. F. Shan, W. Yang, X. G. Tang, M. B. Yang, B. H. Xie, Q. Fu and Y. W. Mai, *Polym. Test.*, 2010, **29**, 273–280.
- 54 C. H. Kung, P. K. Sow, B. Zahiri and W. Mérida, *Adv. Mater. Interfaces*, 2019, **6**, 1900839.
- 55 N. Verplanck, Y. Coffinier, V. Thomy and R. Boukherroub, *Nanoscale Res. Lett.*, 2007, **2**, 577–596.
- 56 F. Zhang and H. Y. Low, *Langmuir*, 2007, **23**, 7793–7798.
- 57 H. Gu, C. Wang, S. Gong, Y. Mei, H. Li and W. Ma, *Surf. Coat. Technol.*, 2016, **292**, 72–77.
- 58 R. Belibel, T. Avramoglou, A. Garcia, C. Barbaud and L. Mora, *Mater. Sci. Eng. Carbon*, 2016, **59**, 998–1006.
- 59 M. Safari, A. Mugica, M. Zubitur, A. M. de Ilarduya, S. Muñoz-Guerra and A. J. Müller, *Polymers*, 2020, **12**, 17.
- 60 M. Safari, I. Otaegi, N. Aramburu, G. Guerrica-Echevarria, A. M. De Ilarduya, H. Sardon and A. J. Müller, *Polymers*, 2021, **13**, 2263.
- 61 C. M. Chan and L. T. Weng, *Materials*, 2016, **9**, 655.
- 62 K. L. A. Cimatú, U. I. Premadasa, T. D. Ambagaspiitiya, N. M. Adhikari and J. H. Jang, *J. Colloid Interface Sci.*, 2020, **580**, 645–659.
- 63 G. Zorn, F. I. Simonovsky, B. D. Ratner and D. G. Castner, *Adv. Healthcare Mater.*, 2022, **11**, 2100894.
- 64 Y. Liu, E. Ranucci, M. S. Lindblad and A. C. Albertsson, *J. Polym. Sci., Part A:Polym. Chem.*, 2001, **39**, 2508–2519.
- 65 L. Diao, K. Su, Z. Li and C. Ding, *RSC Adv.*, 2016, **6**, 27632–27639.
- 66 Y. Tsai, C. H. Fan, C. Y. Hung and F. J. Tsai, *Eur. Polym. J.*, 2009, **45**, 115–122.
- 67 Y. L. Li, X. M. Jia, X. Z. Zhang, Z. Y. Lu and H. J. Qian, *Soft Matter*, 2022, **19**, 128–136.
- 68 S. Feng, H. Yang and Z. Qiu, *Polym. Chem.*, 2025, **16**, 2133–2142.
- 69 Y. Zhuang, X. Liu and Y. Gu, *Polym. Chem.*, 2012, **3**, 1517–1525.
- 70 H. Hu, R. Zhang, W. Bin Ying, L. Shi, C. Yao, Z. Kong, K. Wang, J. Wang and J. Zhu, *Polym. Chem.*, 2019, **10**, 1812–1822.
- 71 K. S. Savitha, B. R. Paghadar, M. Senthil Kumar and R. L. Jagadish, *Polym. Chem.*, 2022, **13**, 3562–3612.

