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Impact of hydrolysis pretreatment on the compostability of biodegradable poly(caprolactone) and poly(lactic acid) films†

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The biodegradation performance of non-pretreated and pretreated commercial polyesters was evaluated under simulated composting conditions to understand how abiotic pretreatment accelerates biotic degradation. Polylactic acid (PLA) and polycaprolactone (PCL) were subjected to hydrolysis pretreatment and assessed under simulated composting conditions for 120 days. In addition to tracking CO₂ evolution, polymer-intrinsic factors such as chain scission, measured by reductions in intrinsic viscosity molecular weight (M_{η}), and changes in crystallinity (X_c) were also evaluated for both non-pretreated and pretreated samples during the biodegradation process. Hydrolysis pretreatment resulted in a reduction of initial M_{η} and an increase of initial X_c for all polymer samples. The initial decrease in M_{η} was particularly marked for PLA, which showed about 30% decrease, while PCL exhibited a reduction of just around 7%. Regarding initial X_c , the most significant increase was also seen in PLA, which jumped from approximately 0% to c. 30%. Hydrolysis of semi-crystalline polymers primarily affects the amorphous region, where elevated temperatures allow water to break polymer chains easily. However, for PLA, the disruption of the crystalline structure leads to a less stable type of crystal, probably due to an increase in the rigid amorphous region that enhances the overall biodegradation process. The effect of pretreatment on the biotic phase showed minimal differences for PCL but a noticeable overall increase in biodegradation for the pretreated PLA.

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1. Introduction

Recent legislative developments in biotechnology and biomanufacturing highlight the need for increased bio-based and recyclable-by-design polymers capable of replacing over 90% of current plastics and commercial polymers at scale by 2030.¹ Initially, this initiative only partially considered managing biodegradable materials due to their lower production levels than traditional non-biodegradable polymers. However, industries have been shifting towards replacing single-use plastics with reusable, recyclable, and even industrial and home-compostable options to foster sustainable practices. Nonetheless, recycling infrastructure has struggled to meet ambitious corporate recycling goals, as approximately 25 to 47% of collected materials are lost in today's recycling processes, underscoring the necessity to enhance product recyclability and improve recycling rates.² Organic waste currently comprises approxi-

mately one-third of municipal solid waste globally.³ Implementing compostable plastics could provide a viable and environmental solution to reduce plastic waste accumulation in the environment while also decreasing the amount of plastic waste sent to landfills and incinerators.⁴ Moreover, biodegradable and compostable materials could significantly enhance the end-of-life (EoL) management of organic waste, especially in cases of food-contaminated packaging—such as packaging for meat, microwaveable frozen foods, and soft cheeses—which typically cannot be recycled due to contamination.⁵ Instead, these materials could effectively be diverted to composting facilities. Current research, market trends, and consumer preferences collectively indicate a growing demand for sustainable, biodegradable alternatives, particularly for single-use packaging applications.

The transition to biodegradable polymers is a critical step in reducing plastic waste; however, challenges persist in ensuring effective degradation and in avoiding unintended environmental consequences, such as microplastic formation. Microplastics, commonly found in paint, tires, pellets, and personal care items, pose significant environmental risks. Microplastic emissions are projected to rise from 9 million tonnes in 2019 to 16 million tonnes by 2040, and existing measures to prevent their release remain insufficient.⁶ The variety of sources generating microplastics from macro plastics

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Hydrolysis emerges as an effective and practical pretreatment method that addresses many of the challenges other pre-

PCL and PLA, both polymers, are capped with a carboxylic acid terminal group. When examining the effect of hydrolysis on PLA in an aqueous solution, a self-catalytic hydrolysis reaction is observed, which is not seen with PCL.¹² Consequently, the hydrolysis kinetics differ between PCL and PLA, as PLA undergoes a self-catalyzed reaction. Additionally, PCL's faster crystallization rate compared to PLA results in greater crystallinity in the materials observed after degradation. Furthermore,

treatment techniques present. It can break down the amorphous regions of a polymer to increase the number of available sites for enzymatic attack, the biotic step initiated by microorganisms.²² Hydrolysis also applies to real-life scenarios, as polymers like PLA and PCL naturally undergo hydrolysis when exposed to moisture, leading to gradual degradation over time. However, specific temperature conditions are crucial for gaining meaningful insights into the hydrolysis process.¹⁹ While previous studies have utilized enzymatic hydrolysis, some have struggled to determine whether the weight loss of the polymer resulted from complete biodegradation or from the hydrolytic breakdown of chains into low-molar-mass segments.²⁶ This highlights the need for research that measures M_n both after hydrolysis pretreatment and during biodegradation to assess where weight loss occurs accurately. Hydrolysis is less expensive than other pretreatment methods and offers easy accessibility regarding the necessary components.²²

Since hydrolysis primarily affects the amorphous regions of each polymer, a hydrolysis pretreatment is likely to have a more significant impact on PLA than on PCL regarding final biodegradation. Therefore, PLA and PCL are ideal candidates for comparing how hydrolysis pretreatment influences their M_n and X_c , directly affecting their biodegradation rates. Their distinct physiochemical properties and similar applications in biodegradable polymers make them particularly suitable for this comparison. PLA is recognized for its high rate of hydrolyzed degradation due to its autocatalytic properties, establishing it as a common benchmark in biodegradation studies alongside other biodegradable polymers.^{12,28} Meanwhile, PCL is a preferred biodegradable polymer because of its versatile mechanical properties and the capacity to adjust degradation kinetics by varying factors such as X_c , M_n , and structural porosity (e.g., porosity and thickness).¹⁸

This work aimed to evaluate the biodegradation performance under composting conditions of two aliphatic biodegradable polyesters, biobased PLA and fossil-based PCL, both pretreated by hydrolysis. In addition to tracking CO_2 evolution, polymer-intrinsic factors such as chain scission, measured by M_n reduction, and X_c evolution were also assessed for both non-pretreated and pretreated samples during the biodegradation process.

2. Materials and methods

Ingeo™ biopolymer 2003D poly (96% L-lactic acid) (PLA) resin was purchased from NatureWorks LLC (Minnetonka, MN, US), with a number (M_n) and weight average molecular weight (M_w) previously reported in the range of 120–121 kDa and 200–235 kDa, respectively.²⁹ PCL CAPA 6800D with an M_w reported by the manufacturer of 80 kDa was obtained from Ingevity® (North Charleston, SC, US). Cellulose powder (particle size *c.* 20 μm) and chloroform Omnisolv were obtained from Sigma Aldrich (Milwaukee, WI, US).

2.1. Preparation of samples

The PLA and PCL samples were produced using compression molding. Each sheet was made from five grams of either PLA or PCL pellets. Before compressing the PLA or PCL pellets, aluminum foil sheets were preheated for 10 minutes. The PLA pellets sandwiched between two aluminum foils were compressed at 180 °C for three minutes at 33 MPa in a hot press (PHI, City of Industry, CA, US). Meanwhile, the PCL pellets were compressed at 90 °C for three minutes at 33 MPa in the hot press. Immediately after hot pressing, the produced sheets were quenched by storing them in a container with ice dry (*c.* −79 °C) for 10 minutes to prevent recrystallization. After production, all samples were kept at −15 °C until use. The average thickness of the PLA samples was $184 \pm 84 \mu\text{m}$ ($7.2 \pm 3.3 \text{ mil}$), while the average thickness of the PCL samples was $193 \pm 54 \mu\text{m}$ ($7.6 \pm 2.1 \text{ mil}$), both determined by averaging ten measurements for each sample using a digital micrometer (Testing Machines Inc., Ronkonkoma, NY, US).

2.2. Pretreatment by hydrolysis

The hydrolysis test was conducted using a batch immersion method in HPLC-grade water under unbuffered conditions. Briefly, PLA and PCL films measuring $1 \times 1 \text{ cm}^2$ were introduced into deionized water at a ratio of approximately 0.62 mL cm^{-2} for PLA and 0.65 mL cm^{-2} for PCL. The hydrolysis conditions for PLA were set at 95 °C for 5 hours, while the PCL samples underwent hydrolysis at 45 °C for 72 hours. After retrieval, the samples were thoroughly dried before further testing and analysis.

2.3. Viscosity-average molecular weight (M_η) determination

To calculate the M_η , the Mark–Houwink equation was employed using the intrinsic viscosity (η) obtained from measurements conducted in accordance with ASTM D2857-16, using chloroform (CHCl_3) as the solvent. The intrinsic viscosity was determined using a Rheotek™ RPV-1 (PSL Rheotek, IN, US) automated viscosity measuring instrument, with a Rheotek™ iSP-1 preparation system operating at 30 ± 0.1 °C for PLA and PCL. Each sample was prepared in four different concentrations ranging from 0.125 to 0.500 g dL^{-1} . The reduced viscosity and inherent viscosity were plotted against concentration. The intrinsic viscosity value was derived by extrapolating the reduced viscosity and inherent viscosity to zero concentration. The average of the two obtained intercept values was used as the intrinsic viscosity to determine the M_η using the Mark–Houwink equation:

$$[\eta] = K \cdot M_\eta^\alpha \quad (1)$$

where K and α are constants, which depend on the nature of the polymer, solvent, and temperature obtained from the intercept and slope of a double logarithmic plot of intrinsic viscosity *versus* M_η . The K and α values of each material in CHCl_3 at 30 °C were obtained from the literature and shown in Table 1.



Table 1 K and α values for PLA and PCL for the Mark–Houwink equation

Polymer	Temperature, °C	K , 10^{-3} mL g $^{-1}$	α	Ref.
PLA	30	0.0131	0.777	30
PCL	30	0.0130	0.828	31

2.4. Crystallinity (X_c) measurement

The X_c , glass transition temperature (T_g), crystallization temperature (T_c), and melting temperature (T_m) of PLA and PCL were measured using a Differential Scanning Calorimetry (DSC) model Q100 (TA Instruments, New Castle, DE, US) equipped with a mechanical cooling system performed under the heat-only condition with a heating rate of 10 °C min $^{-1}$ from 0 to 180 °C. The degree of X_c was determined using eqn (2) below:

$$X_c = \frac{\Delta H_m - \sum \Delta H_c}{\Delta H_m^0} \quad (2)$$

where ΔH_m is the enthalpy of fusion from the integration of the heat flow in the melting region, $\sum \Delta H_c$ is the sum of the exothermic enthalpy peaks, and ΔH_m^0 is the calculated melting enthalpy of 100% crystalline polymers obtained from the literature. For PCL, this value is 167 J g $^{-1}$ (ref. 32) and for PLA is 93 J g $^{-1}$.³³

2.5. Biodegradation test

PLA and PCL films (8 g) were cut into ~1 cm 2 pieces and introduced into a bioreactor (1.9 L) filled with mixed manure-straw compost that had been screened with 10 mm mesh and acquired from the Michigan State University (MSU) Composting Facility (East Lansing, MI, US), along with vermiculite equilibrated with deionized water to enhance the moisture content of the compost (up to 50%) in a 4 : 1 ratio (based on dry weight compost). The measurement of CO $_2$ evolution was conducted for PLA and PCL films in triplicate bioreactors, and the average and standard error values are reported. One additional bioreactor was used for sampling to study M_n reduction. Three blank bioreactors (compost without samples) were employed to detect the background CO $_2$ evolution. Three bioreactors with 8 g of cellulose powder were included as a positive control due to cellulose's high biodegradability. The characterization of the compost's physicochemical parameters was conducted by the MSU Soil and Plant Nutrient Laboratory (East Lansing, MI, US), and the results are presented in Table S1 in the ESI.† The biodegradability was evaluated in compost under aerobic conditions using a direct measurement respirometer system, where temperature (58 ± 2 °C), relative humidity ($50 \pm 5\%$), and air flow rate (40 ± 2 cm 3 min $^{-1}$) were the primary controlled parameters. The methodology and equipment are detailed thoroughly elsewhere.³⁴ The CO $_2$ evolved from the blank bioreactor was considered the background signal, and this value was subtracted from the amount of CO $_2$ produced by each sample bioreactor to calculate the biodegradation of each sample; where the percentage bio-

degradation represents the total amount of carbon molecules converted to CO $_2$, calculated according to eqn (3) below:

$$\% \text{ of Biodegradation} = \frac{(\text{CO}_2)_t - (\text{CO}_2)_b}{M_t \times C_t \times \frac{44}{12}} \times 100 \quad (3)$$

where the numerator is the difference between the average of the three bioreactors' cumulative mass of CO $_2$ evolved for the sample (CO $_2$) $_t$ and the average CO $_2$ evolved from the three blank bioreactors (CO $_2$) $_b$. The denominator represents the theoretical amount of CO $_2$ produced by the sample. M_t is the total mass of the sample, C_t is the proportion of carbon present in that sample as determined by CHN analysis, and 44 and 12 are the molecular mass of CO $_2$ and the atomic mass of carbon, respectively.³⁵ The carbon content for each sample was quantified using a PerkinElmer 2400 Series II CHNS/O Elemental Analyzer (Shelton, CT, US), and results can be found in Table S2 in the ESI.†

3. Results and discussion

The leading factors that affect polymer biodegradation are related to the polymer's structure and the environment. Pretreatment focuses on deteriorating the intrinsic polymer structure, such as polymer chain scission (observed by a reduction in M_n) and crystallinity structure. We have previously reported that abiotic pretreatment of aliphatic polyesters could be reported as a feasible way to enhance overall biodegradation performance, particularly under composting conditions.²² Targeting the reduction of M_n has been one of these methods. In this study, the reduction of M_n was achieved through hydrolysis at high temperatures (above the T_g of the samples), and the results of biodegradation performance comparisons between non-pretreated and pretreated samples are discussed.

3.1. Effect of abiotic pretreatment on initial molecular weight and initial crystallinity

During hydrolysis pretreatment, water penetrates the entire polymer, causing random hydrolytic chain splits to occur uniformly throughout the matrix. The ester chains are cleaved, forming carboxyl terminal groups that decrease the size of the molecular chains, leading to a reduction in M_n .³⁶ Previous studies have shown that M_n of polymers declines with increased exposure time in water during hydrolysis.¹⁹ Table 2 displays the M_n of PCL and PLA before and after hydrolysis. The M_n of PLA-PT is *c.* 130 kDa, compared to *c.* 167 kDa for the non-pretreated sample, indicating a 22% reduction. Similarly, for PCL-PT, the M_n is *c.* 93.6 kDa, whereas the M_n of the non-pretreated PCL is *c.* 97 kDa, showing a 3.5% reduction. Apart from crystallinity, the high hydrophobicity of PCL has been recognized as a crucial factor for the slow decrease of M_n .³⁷ This underscores the importance of considering surface properties and their alteration through acidity/alkaline treatment to expedite hydrolysis, enzymatic depolymerization, and the overall degradation process in the case of polyesters.³⁸



3.2.2. Evolution of crystallinity and reduction of molecular weight. Fig. 3 shows the evolution of X_c , M_n reduction, CO_2 evolution, and % biodegradation. The evolution of X_c was measured during the first 30 days of the biodegradation test for both PLA and PCL samples when samples could be collected. Throughout this initial biodegradation period, X_c increased for all samples, with PCL and PCL-PT converging at similar X_c values around day 22, and PLA and PLA-PT approaching similar X_c values around day 28 (Fig. 3a and b).

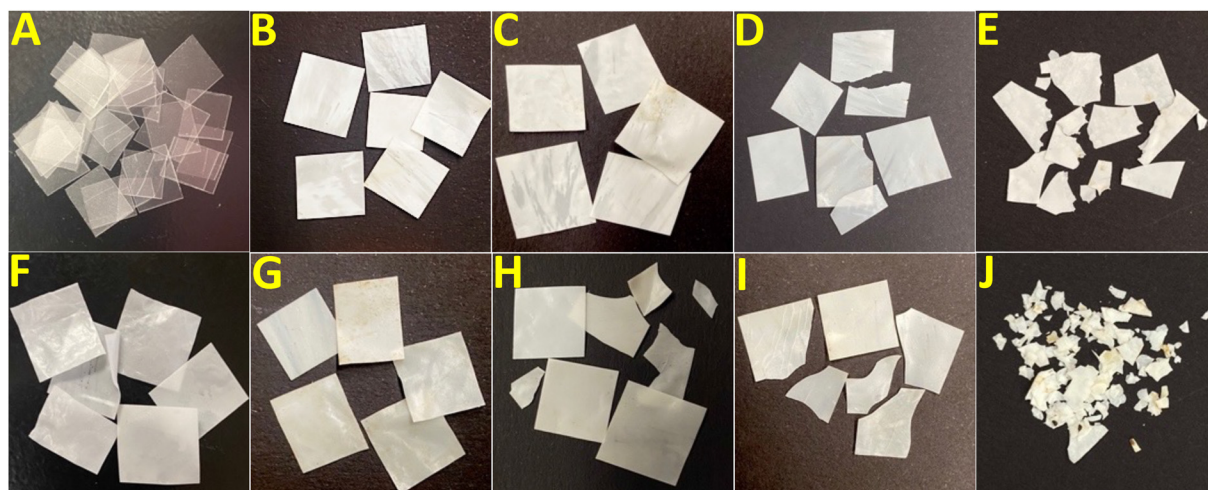


Fig. 2 PLA non-pretreated day 0 (A), day 6 (B), day 12 (C), day 18 (D), day 30 (E) and PLA pretreated day 0 (F), day 6 (G), day 12 (H), day 18 (I), day 30 (J) in compost. Note: Initial samples were cut to an area of $\sim 1 \text{ cm}^2$.

Crystallinity increases during biodegradation as hydrolysis primarily breaks down amorphous regions, leaving the more resistant crystalline structures intact. As polymer chains degrade and shorten, they gain mobility, allowing the amorphous regions to reorganize into crystalline structures.²⁸ Studies on PCL demonstrated that amorphous regions degrade first, leaving behind crystalline material that eventually breaks down. Both PLA and PCL undergo hydrolytic degradation through bulk erosion, where the entire material degrades simultaneously.³⁹ This selective bulk erosion mechanism of degradation enhances overall crystallinity and slows further degradation, as crystalline regions resist hydrolysis and microbial attack, thereby prolonging material longevity. However, it has been reported that surface erosion patterns, such as grooves and cracks observed in PCL exposed to fungi in compost, further illustrate the persistence and evolution of crystalline structures during biodegradation.⁴⁰ Furthermore, Tsuji and Suzuyoshi found that the environmental degradation of PCL can also occur *via* surface erosion rather than bulk erosion, with degradation happening inhomogeneously on the film surface due to microbial attachment.⁴¹ Considering the critical thickness model theory, PCL and PCL-PT samples are deemed to degrade following a bulk erosion mechanism in our test.^{39,42} Critical sample thickness for PCL has been reported with a value of *c.* 1.3 cm, indicating a preferred bulk erosion mechanism below that theoretical value.⁴³ PLA and PLA-PT samples also align with this model, as the critical thickness for poly(α -hydroxy esters) is reported as *c.* 7.4 cm. In this study, the tested samples' thickness remained below this threshold, supporting the conclusion that PLA and PLA-PT samples degraded through a bulk erosion mechanism.³⁹

The M_n was measured for samples retrieved during the biodegradation test for as long as the samples were available. The values of M_n for PLA-PT could not be measured on day 30 using the viscosity technique. In the case of PCL, samples were collected until day 18 of the biodegradation test. Tables S3 and

S4 in the ESI† present the M_n reduction for PLA and PLA-PT, as well as for PCL and PCL-PT, respectively, while Fig. 3c and d illustrate the normalized M_n . PLA showed an 86% reduction in M_n over the 30 days in compost, while PLA-PT experienced a 70% decrease between day 0 and day 18. For PCL, the M_n decreased by 42%, with PCL-PT exhibiting a comparable reduction of 44% during the same 18 day period. The pretreatment performed well for PLA, where a significant portion of the amorphous region was hydrolyzed, achieving destabilization of the three-phase model structure (*i.e.*, crystalline, and rigid and mobile amorphous regions). This allowed for a rapid reduction in M_n during the first month of the biodegradation process at a temperature around the T_g of PLA. The high initial X_c influenced the overall decrease in M_n for PCL, displaying similar final values before and after pretreatment.

3.2.3. Evolution of CO_2 and biodegradation. In the case of PLA, the pretreatment resulted in an effective reduction of the abiotic phase by about a week compared to PLA non-pretreated, accelerating the biotic phase. The high rate of PLA-PT biotic phase reached biodegradation of *c.* 100% at day 50, while non-pretreated PLA biodegradation was *c.* 75% at the same point (Fig. 3f and h). In the case of PCL, a small advantage of the pretreatment is observed for PCL-PT, reaching *c.* 100% on day 70, while PCL was *c.* 80% on the same day (Fig. 3e and g). The high initial reduction of M_n for PLA (*c.* 22%) played a significant role in reducing the extension of the lag phase and triggering the biotic phase. Furthermore, the increase in initial X_c for PLA by *c.* 30% is indicative that the amorphous phase of PLA structure was highly affected by the hydrolysis pretreatment.

The positive control cellulose reached biodegradation of *c.* 100% at *c.* 40 days. For PLA, abiotic degradation was observed as the controlling mechanism in compost during the first 20 days. A slight acceleration of the abiotic stage was observed for PLA-PT compared to non-pretreated PLA (raw data for biodegradation is provided in Tables S5–S10 in the ESI†).



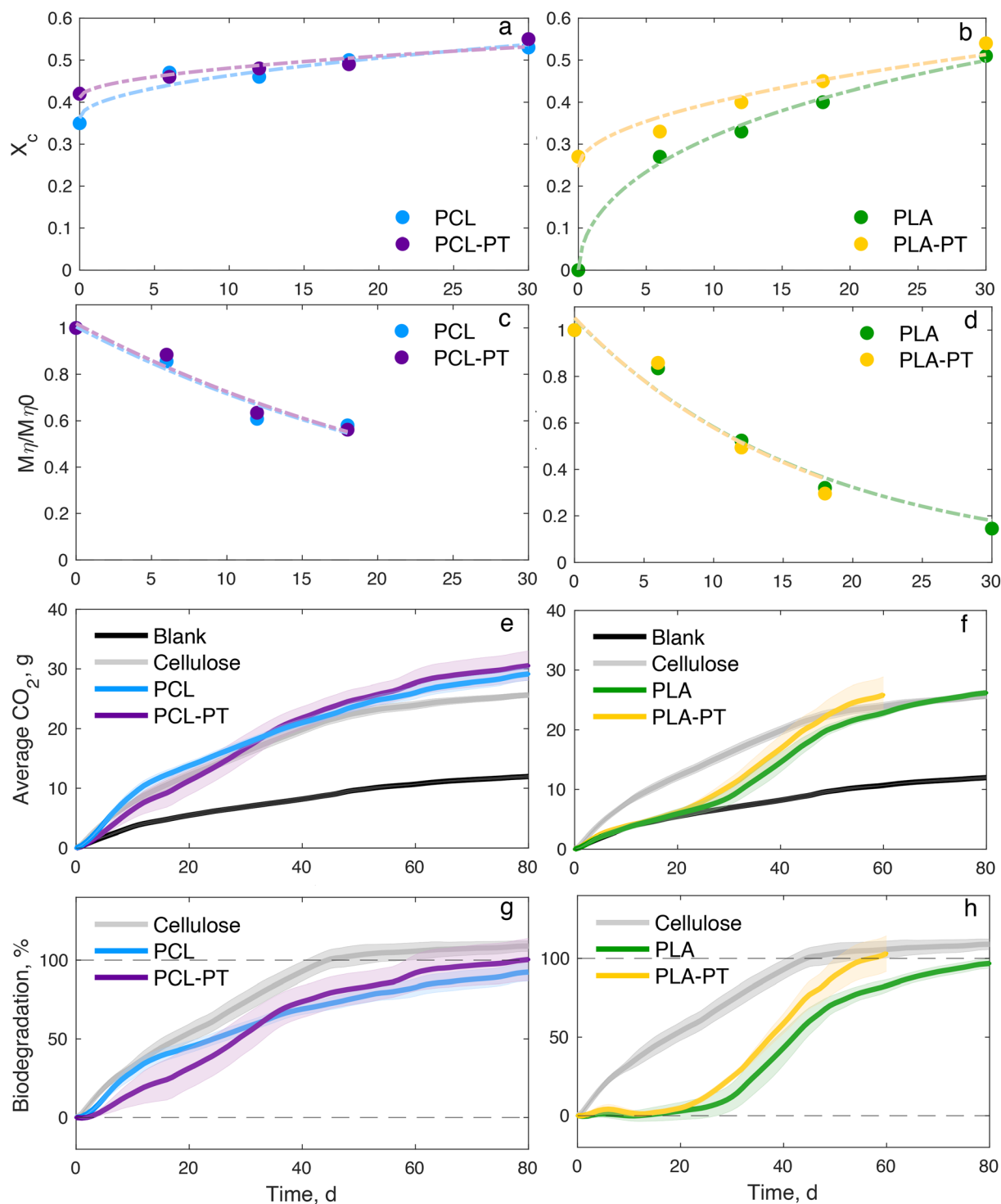


Fig. 3 (a) crystallinity (X_c) evolution of pretreated and non-pretreated PCL, (b) crystallinity (X_c) evolution of pretreated and non-pretreated PLA, (c) molecular weight (M_n) of pretreated and non-pretreated PCL, (d) molecular weight (M_n) of pretreated and non-pretreated PLA, (e) CO_2 evolution of pretreated and non-pretreated PCL, (f) CO_2 evolution of pretreated and non-pretreated PLA, (g) biodegradation of pretreated and non-pretreated PCL, (h) biodegradation of pretreated and non-pretreated PLA until day 80. The shade in the background for each material represents the standard error between replicates.

3.3. Evolution of thermal properties of samples during biodegradation

Fig. 4 presents the thermal analysis for PCL, PCL-PT, PLA, and PLA-PT samples for day 0 and during the first month of degradation. The T_g , T_c , and T_m for PCL are reported in the literature as *c.* -60 °C, 28 °C, and 60 °C, respectively,⁴⁴ while PLA typically falls within 50 – 65 °C for T_g , *c.* 94.5 °C for T_c , and 130 – 180 °C for T_m .^{45,46} For this work, DSC values measured at day 0 are displayed in Table 2. The average value for PCL was around 52 °C for T_m . After pretreatment, T_m increased to *c.* 56.5 °C due to increasing in crystal thickness and perfection indicates higher stability of the crystals.⁴⁷ The low temperature melting shoulders were not observed for PCL in the thermophilic samples. This may be attributed to crystal rearrangements occurring at higher temperatures under thermophilic conditions.⁴⁰ Untreated PLA at day 0 exhibited T_g and T_m values of around 55.5 °C and 144 °C, respectively. PLA-PT showed a T_g of 60.3 °C and T_m of 148.8 °C. A significant shift toward higher temperatures for the T_m is observed for PCL compared to PCL at day 0, indicating a more ordered and crystalline structure during early stage of biodegradation. However, for PLA, the T_m shows a shift towards lower temperature indicative of the formation of imperfect crystals, especially for PLA-PT samples due to the decrease of M_n , following the behavior of an adapted Flory–Fox equation reported by Saeidlou *et al.*, 2012 below (4). From the double melting peak displayed can be deducted the formation of α -crystal for the highest T_m and δ -crystal form for the lowest T_m .⁴⁸

$$T_m = T_m^\infty - \frac{A}{M_n} \quad (4)$$

where $T_m^\infty = 181.3$ °C and $A = 1.02 \times 10^5$ °C g mol⁻¹.⁴⁸ During crystallization both forms α - and δ -crystal are formed with a similar energy and, therefore, able to coexist. From this, the higher T_m corresponds to the more stable α form, while the lower T_m is associated with the less stable δ -crystal.⁴⁹ The shift to a lower temperature for T_m of PLA-PT is related to an increase in the crystallinity but indicative of a least ordered crystalline structure probably associated with a δ -structure formation at day 30. The T_m of δ structure has been reported to be *c.* 10 °C lower than α -crystal, suggesting a less stable δ form.⁴⁸

3.4. Relationship between crystallinity and molecular weight reduction during biodegradation

Abiotic pretreatment may provide advantages over biotic pretreatment by enabling quicker disintegration and increasing the exposed area for microbial attack, while also reducing the M_n of the structure and total disintegration of the amorphous region. Additionally, a more resilient X_c typically diminishes abiotic hydrolysis, enzymatic hydrolysis, and the overall biodegradation process. Furthermore, the behavior of surface properties must be considered when enhancing biodegradation performance.

During bulk degradation, when water diffusion into the polymer occurs faster than the hydrolytic attack, the M_n decreases early in the process. This decrease in M_n , along with the increased mobility of shorter polymer segments, the presence of water acting as a plasticizer, and elevated temperatures, can lead to a higher degree of X_c .⁵⁰ The increase in initial X_c due to the reduction of initial M_n is particularly relevant for PLA, but it has minimal effect on PCL due to its larger

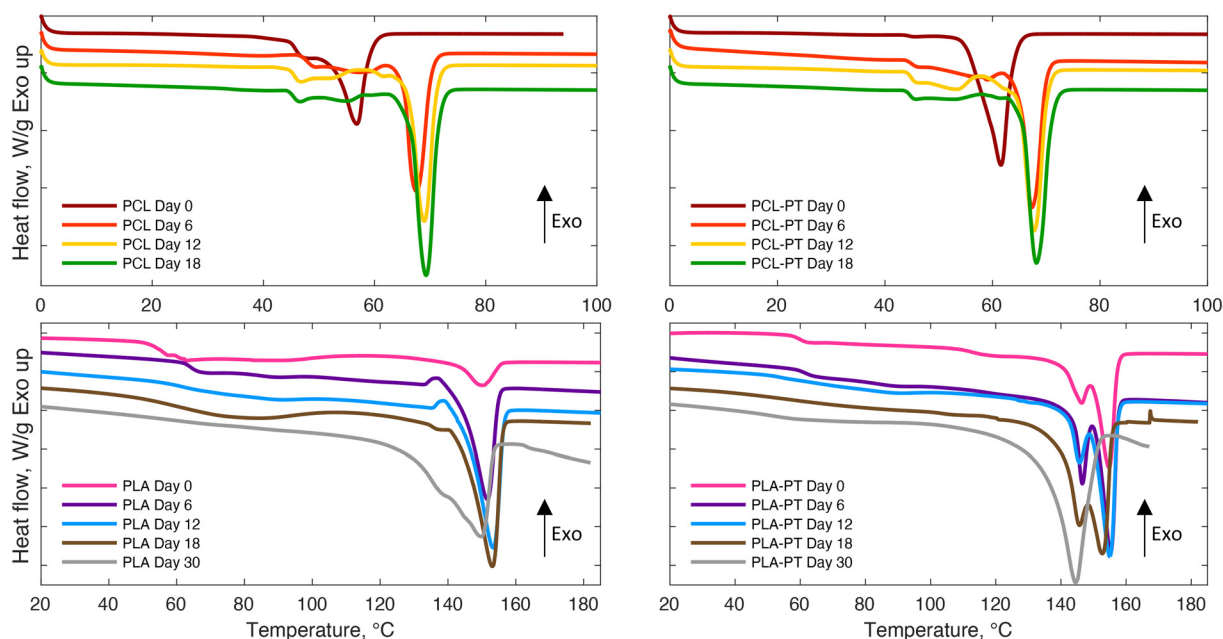


Fig. 4 Differential scanning calorimetry of PCL (top left), PCL pretreated (top right), PLA (bottom left), and PLA pretreated (bottom right) for day 0 and during the early stage of the biodegradation process.



initial X_c . In the case of PLA, the impact of M_n reduction on fragmentation and the initiation of depolymerization is more pronounced than for PCL.

For PLA, hydrolysis pretreatment lowers T_g and decreases PLA's M_n due to chain scission, enhancing chain mobility and increasing the biodegradation rate. However, the increased X_c formed during hydrolysis can initially slow degradation since crystalline regions are tightly packed and less accessible to water and microorganisms. In contrast, PCL, which starts with a high degree of crystallinity, experiences minimal changes in both M_n and X_c after hydrolysis. Consequently, the biodegradation rate of PCL remains relatively unaffected, as its well-formed crystalline structure does not undergo significant changes during hydrolysis.¹⁸

A common characteristic of hydrolytic degradation processes is that increasing X_c tends to lower the degradation rate. This occurs because water cannot easily penetrate the highly ordered crystalline regions.³⁹ Although fast crystallization may initially slow biodegradation due to the resistance of crystalline regions, the interaction between crystalline and amorphous regions complicates the prediction. In the rigid amorphous regions (RAFs) formed between the mobile amorphous regions (MAF) and the crystalline domains, the degradation rate could be higher, resulting in an increased biodegradation rate and a more nuanced effect on overall biodegradation,⁵¹ which remains to be decoupled.

4. Conclusions

Using hydrolysis as a green and affordable alternative to accelerate biodegradation showed significant improvements for PLA but only modest gains for PCL. This is indicated by the fact that the effect of pretreatment was more pronounced in reducing M_n for PLA than for PCL, thus reducing the lag phase of pretreated PLA. When possible, it is essential to ascertain the best optimization strategies for pretreatment, considering the general characteristics of biodegradable polymers, especially polyesters, while also taking into account critical parameters of the media, such as pH for hydrolysis pretreatments at high temperatures. Additionally, altering the surface properties of polymer films through pretreatment can be beneficial, leading to a more effective breakdown process. Understanding the unique attributes of each polymer will facilitate more efficient pretreatments and biodegradation strategies.

Author contributions

Jordan D'Amario: Experimental run, data collection, writing – original draft, editing, and reviewing, visualization. Anibal Bher: Conceptualization, experimental run, data collection and curation, writing – original draft, editing, and reviewing, supervision, validation. Wanwarang Limsukon: Experimental run, data collection and curation, visualization, writing – original

draft, editing, and reviewing. Rafael Auras: conceptualization, writing – original draft, editing and reviewing, funding acquisition, supervision. All the authors approved the final version of the manuscript.

Data availability

Data is available in the manuscript, and additional raw data supporting this article has been included as part of the ESI.†

Conflicts of interest

There are no conflicts to declare.

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References

- 1 The White House, *Bold Goals for U.S. Biotechnology and Biomufacturing: Harnessing Research and Development to Further Societal Goals*, 2023.
- 2 UNEP, *Turning off the Tap How the world can end plastic pollution and create a circular economy*, 2023, Available from: <https://www.unep.org/resources/turning-off-tap-end-plastic-pollution-create-circular-economy>.
- 3 World Bank, *What a Waste 2.0 A Global Snapshot of Solid Waste Management to 2050 Overview*, 2018.
- 4 X. E. Crystal Thew, S. C. Lo, R. N. Ramanan, B. T. Tey, N. D. Huy and O. Chien Wei, Enhancing plastic biodegradation process: strategies and opportunities, *Crit. Rev. Biotechnol.*, 2024, **44**(3), 477–494.
- 5 Closed Loop Partners, *The Realities of Compostable Packaging Disintegration in Composting Systems Breaking it Down*, 2024, Available from: <https://www.closedlooppartners.com>.
- 6 Nordic Council of Ministers, *Towards Ending Plastic Pollution by 2040*, 15 Global Policy Interventions for Systems Change, 2023, Available from: <https://www.norden.org>.
- 7 R. C. Thompson, W. Courteney-Jones, J. Boucher, S. Pahl, K. Raubenheimer and A. A. Koelmans, Twenty years of microplastics pollution research—what have we learned?, *Science*, 2024, **386**(6720), eadl2746, Available from: <https://www.science.org/doi/10.1126/science.adl2746>.



- 8 S. Nizamuddin, A. J. Baloch, C. Chen, M. Arif and N. M. Mubarak, Bio-based plastics, biodegradable plastics, and compostable plastics: biodegradation mechanism, biodegradability standards and environmental stratagem, *Int. Biodeterior. Biodegrad.*, 2024, **195**, 105887.
- 9 R. Auras, H. Tsuji, S. Selke and L. Loong-Tak, *Poly(Lactic Acid) synthesis, structures, properties, processing*, Wiley-Blackwell, 2022.
- 10 L. K. Ncube, A. U. Ude, E. N. Ogunmuyiwa, R. Zulkifli and I. N. Beas, Environmental impact of food packaging materials: A review of contemporary development from conventional plastics to polylactic acid based materials, *Materials*, 2020, **13**, 1–24.
- 11 C. Sun, S. Wei, H. Tan, Y. Huang and Y. Zhang, Progress in upcycling polylactic acid waste as an alternative carbon source: A review, *Chem. Eng. J.*, 2022, **446**, 136881.
- 12 G. L. Siparsky, K. J. Voorhees and F. Miao, Hydrolysis of Polylactic Acid (PLA) and Polycaprolactone (PCL) in Aqueous Acetonitrile Solutions: Autocatalysis, *J. Environ. Polym. Degrad.*, 1998, **6**, 31–41.
- 13 M. A. Woodruff and D. W. Hutmacher, The return of a forgotten polymer - Polycaprolactone in the 21st century, *Prog. Polym. Sci.*, 2010, **35**, 1217–1256.
- 14 M. Thakur, I. Majid, S. Hussain and V. Nanda, Poly(ϵ -caprolactone): A potential polymer for biodegradable food packaging applications, *Packag. Technol. Sci.*, 2021, **34**, 449–461.
- 15 A. Bher, I. U. Unalan, R. Auras, M. Rubino and C. E. Schvezov, Graphene modifies the biodegradation of poly(lactic acid)-thermoplastic cassava starch reactive blend films, *Polym. Degrad. Stab.*, 2019, **164**, 187–197.
- 16 H. Wan, C. Sun, C. Xu, B. Wang, Y. Chen, Y. Yang, *et al.*, Synergistic reinforcement of polylactic acid/wood fiber composites by cellulase and reactive extrusion, *J. Cleaner Prod.*, 2024, **434**, 140207.
- 17 S. S. Hamdani, H. M. Elkholy, M. O. Alghaysh, I. Wyman, A. Bher, R. Auras, *et al.*, Recyclable and Biodegradable Paper Coating with Functionalized PLA and PBAT, *ACS Omega*, 2025, **10**(11), 11483–11497.
- 18 I. Castilla-Cortázar, J. Más-Estells, J. M. Meseguer-Dueñas, J. L. Escobar Ivirico, B. Marí and A. Vidaurre, Hydrolytic and enzymatic degradation of a poly(ϵ -caprolactone) network, *Polym. Degrad. Stab.*, 2012, **97**(8), 1241–1248.
- 19 W. Limsukon, M. Rubino, M. Rabnawaz, L. T. Lim and R. Auras, Hydrolytic degradation of poly(lactic acid): Unraveling correlations between temperature and the three phase structures, *Polym. Degrad. Stab.*, 2023, **217**, 110537.
- 20 Waste Dive, *Compostable packaging's relationship status with organics recyclers: 'It's complicated'*, 2024, Available from: <https://www.wastedive.com/news/bpi-summit-compostable-packaging-usda-calrecycle/> [cited 2024 Sep 25].
- 21 E. Gastaldi, F. Buendia, P. Greuet, Z. Benbrahim Bouchou, A. Benihya, G. Cesar, *et al.*, Degradation and environmental assessment of compostable packaging mixed with biowaste in full-scale industrial composting conditions, *Bioresour. Technol.*, 2024, **400**, 130670.
- 22 A. Bher, Y. Cho and R. Auras, Boosting Degradation of Biodegradable Polymers, *Macromol. Rapid Commun.*, 2023, **44**, e2200769.
- 23 T. Kijchavengkul, R. Auras, M. Rubino, M. Ngouajio and R. T. Fernandez, Assessment of aliphatic–aromatic copolyester biodegradable mulch films. Part I: Field study, *Chemosphere*, 2008, **71**(5), 942–953.
- 24 T. Kijchavengkul, R. Auras, M. Rubino, M. Ngouajio and R. T. Fernandez, Assessment of aliphatic–aromatic copolyester biodegradable mulch films. Part II: Laboratory simulated conditions, *Chemosphere*, 2008, **71**(9), 1607–1616.
- 25 N. Mat Yasin, S. Akkermans and J. F. M. Van Impe, Enhancing the biodegradation of (bio)plastic through pre-treatments: A critical review, *Waste Manage.*, 2022, **150**, 1–12.
- 26 Z. H. W. Easton, M. A. J. Essink, L. Rodriguez Comas, F. R. Wurm and H. Gojzewski, Acceleration of Biodegradation Using Polymer Blends and Composites, *Macromol. Chem. Phys.*, 2023, **224**, 2200421.
- 27 M. Yusefi, M. Khalid, F. M. Yasin, L. C. Abdullah, M. R. Ketabchi and R. Walvekar, Performance of Cow Dung Reinforced Biodegradable Poly(Lactic Acid) Biocomposites for Structural Applications, *J. Polym. Environ.*, 2018, **26**(2), 474–486.
- 28 H. Tsuji, Hydrolytic Degradation, in *Poly(Lactic Acid)*, 2022, pp. 467–516, DOI: [10.1002/9781119767480.ch21](https://doi.org/10.1002/9781119767480.ch21).
- 29 F. Iñiguez-Franco, R. Auras, G. Burgess, D. Holmes, X. Fang, M. Rubino, *et al.*, Concurrent solvent induced crystallization and hydrolytic degradation of PLA by water-ethanol solutions, *Polymer*, 2016, **99**, 315–323.
- 30 J. R. Dorgan, J. Janzen, D. M. Knauss, S. B. Hait, B. R. Limoges and M. H. Hutchinson, Fundamental solution and single-chain properties of polylactides, *J. Polym. Sci., Part B: Polym. Phys.*, 2005, **43**(21), 3100–3111.
- 31 H. Sun, L. Mei, C. Song, X. Cui and P. Wang, The in vivo degradation, absorption and excretion of PCL-based implant, *Biomaterials*, 2006, **27**(9), 1735–1740.
- 32 H. L. Chen, L. J. Li and T. L. Lin, Formation of Segregation Morphology in Crystalline/Amorphous Polymer Blends: Molecular Weight Effect, *Macromolecules*, 1998, **31**(7), 2255–2264, Available from: <https://pubs.acs.org/sharingguidelines>.
- 33 E. W. Fischer, H. J. Sterzel and G. Wegner, Investigation of the structure of solution grown crystals of lactide copolymers by means of chemical reactions, *Z-Polymers*, 1973, **251**, 980–990.
- 34 E. Castro-Aguirre, R. Auras, S. Selke, M. Rubino and T. Marsh, Insights on the aerobic biodegradation of polymers by analysis of evolved carbon dioxide in simulated composting conditions, *Polym. Degrad. Stab.*, 2017, **137**, 251–271, Available from: <https://linkinghub.elsevier.com/retrieve/pii/S0141391017300241>.
- 35 T. Kijchavengkul, R. Auras, M. Rubino, M. Ngouajio and R. Thomas Fernandez, Development of an automatic laboratory-scale respirometric system to measure polymer biodegradability, *Polym. Test.*, 2006, **25**(8), 1006–1016.



- 36 A. Leroux, T. Ngoc Nguyen, A. Rangel, I. Cacciapuoti, D. Duprez, D. G. Castner, *et al.*, Long-term hydrolytic degradation study of polycaprolactone films and fibers grafted with poly(sodium styrene sulfonate): Mechanism study and cell response, *Biointerphases*, 2020, **15**(6), 061006.
- 37 G. G. Pitt, M. M. Gratzl, G. L. Kimmel, J. Surles and A. Sohindler, Aliphatic polyesters II. The degradation of poly (DL-lactide), poly (ϵ -caprolactone), and their copolymers in vivo, *Biomaterials*, 1981, **2**(4), 215–220.
- 38 H. Tsuji, T. Ishida and N. Fukuda, Surface hydrophilicity and enzymatic hydrolyzability of biodegradable polyesters: 1. effects of alkaline treatment, *Polym. Int.*, 2003, **52**(5), 843–852.
- 39 B. Laycock, M. Nikolić, J. M. Colwell, E. Gauthier, P. Halley, S. Bottle, *et al.*, Lifetime prediction of biodegradable polymers, *Prog. Polym. Sci.*, 2017, **71**, 144–189.
- 40 C. Eldsäter, The biodegradation of amorphous and crystalline regions in film-blown poly(ϵ -caprolactone), *Polymer*, 2000, **41**(4), 1297–1304.
- 41 H. Tsuji and K. Suzuyoshi, Environmental degradation of biodegradable polyesters 1. Poly(ϵ -caprolactone), poly[(R)-3-hydroxybutyrate], and poly(L-lactide) films in controlled static seawater, *Polym. Degrad. Stab.*, 2002, **75**(2), 347–355.
- 42 A. Göpferich, Mechanisms of polymer degradation and erosion, *Biomaterials*, 1996, **17**(2), 103–114.
- 43 F. von Burkersroda, L. Schedl and A. Göpferich, Why degradable polymers undergo surface erosion or bulk erosion, *Biomaterials*, 2002, **23**(21), 4221–4231.
- 44 A. Fernández-Tena, R. A. Pérez-Camargo, O. Coulembier, L. Sangroniz, N. Aranburu, G. Guerrica-Echevarria, *et al.*, Effect of Molecular Weight on the Crystallization and Melt Memory of Poly(ϵ -caprolactone) (PCL), *Macromolecules*, 2023, **56**(12), 4602–4620.
- 45 E. Castro-Aguirre, F. Iñiguez-Franco, H. Samsudin, X. Fang and R. Auras, Poly(lactic acid)—Mass production, processing, industrial applications, and end of life, *Adv. Drug Delivery Rev.*, 2016, **107**, 333–366.
- 46 A. Larrañaga and E. Lizundia, A review on the thermo-mechanical properties and biodegradation behaviour of polyesters, *Eur. Polym. J.*, 2019, **121**, 109296.
- 47 A. Wurm, E. Zhuravlev, K. Eckstein, D. Jehnichen, D. Pospiech, R. Androsch, *et al.*, Crystallization and homogeneous nucleation kinetics of poly(ϵ -caprolactone) (PCL) with different molar masses, *Macromolecules*, 2012, **45**(9), 3816–3828.
- 48 S. Saeidlou, M. A. Huneault, H. Li and C. B. Park, Poly(lactic acid) crystallization, *Prog. Polym. Sci.*, 2012, **37**(12), 1657–1677.
- 49 M. Yasuniwa, S. Tsubakihara, K. Iura, Y. Ono, Y. Dan and K. Takahashi, Crystallization behavior of poly(l-lactic acid), *Polymer*, 2006, **47**(21), 7554–7563.
- 50 W. Limsukon, R. Auras and S. Selke, Hydrolytic degradation and lifetime prediction of poly(lactic acid) modified with a multifunctional epoxy-based chain extender, *Polym. Test.*, 2019, **80**, 106108.
- 51 X. Han and J. Pan, A model for simultaneous crystallisation and biodegradation of biodegradable polymers, *Biomaterials*, 2009, **30**(3), 423–430, Available from: <https://linkinghub.elsevier.com/retrieve/pii/S0142961208007424>.

