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Fully bio-based aromatic–aliphatic copolyesters: poly(butylene furandicarboxylate-co-succinate)s obtained by ring opening polymerization†

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Poly(butylene 2,5-furandicarboxylate-co-succinate) copolyesters (coPBF_xS_y) have been synthesized by ring opening polymerization (ROP). Cyclic butylene 2,5-furandicarboxylate and butylene succinate oligomer mixtures, to be used as monomers for ROP, were prepared by high dilution condensation and enzymatic cyclization reactions, respectively. Two different catalytic systems, tin dioctanoate and supported *Candida antarctica* lipase B (CALB), were used for polymerization. Thus two series of copolyesters covering the whole range of compositions were prepared and their properties comparatively examined. In both cases, random copolyesters with compositions close to those used in their respective feeds were obtained. The influence of composition on reaction kinetics with respect to time and temperature was evaluated for the two series. Chemically catalyzed ROP rendered copolyesters with M_w in the ~50 000–65 000 g mol⁻¹ range, whereas values between 15 000 and 45 000 g mol⁻¹ were attained when the ROP reaction was assisted by CALB. The thermal behavior of coPBF_xS_y obtained by ROP was similar to that reported for such copolymers prepared by melt polycondensation. They all start to decompose above 300 °C and display melting enthalpy and temperatures that decrease with copolymerization, attaining minimum values when the comonomer contents are approximate to balance. On the contrary, the glass-transition temperature increased almost linearly with the content of butylene furandicarboxylate units, covering the whole range of values between those of the two parent homopolyesters. Small deviations in thermal properties observed between the two series could be attributed to their differences in molecular weights. Hydrolytic and enzymatic degradation studies revealed that coPBF_xS_y became more degradable with increasing content of succinic units, whereas the homopolyester PBF remained practically unaffected when incubated under similar conditions.

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Introduction

2,5-Furandicarboxylic acid (FDCA) is an aromatic diacid that is industrially prepared by catalytic oxidation of 5-methylfurfural, an aldehyde easily obtained from naturally-occurring hexoses. FDCA stands out as a highly promising bio-based monomer able to replace traditional aromatic diacids of fossil-fuel origin in the production of polyesters and other polycondensates.^{1–3} It is the aromatic nature of the furanic ring that provides enough stiffness to the molecule as to be able to substitute the phthalate units without a deleterious effect on chain mobility. The excellence of FDCA for the preparation of bio-based polyesters and other condensation polymers has been evidenced in

a worthy review recently published by Sousa *et al.*⁴ To date, several poly(alkylene 2,5-furandicarboxylate) homopolyesters have been synthesized and it has been shown that their properties compare well with those displayed by their poly(alkylene terephthalate) analogs. In addition, FDCA, either as the diacid itself or as its dimethyl ester, has been copolymerized with a number of either aromatic or aliphatic diacids to render furanoate-containing copolyesters with satisfactory properties.^{5–10} The most recent progress attained in the production of bio-based 2,5-furandicarboxylate polyesters have been reviewed by Bikiaris *et al.*¹¹

Poly(butylene succinate) (PBS) is a fully bio-based aliphatic polyester that is achieving amazing popularity because it displays good thermal and mechanical behavior in addition to being biodegradable.^{12–14} Copolymerization has been the approach frequently used to render PBS-based materials with higher performance, in particular with improved thermal properties.^{15–18} This is really challenging since renewable com-

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pounds with the appropriate constitution are not easily accessible. Recently, several copolyesters of PBS containing monocyclic or bicyclic carbohydrate units with improved thermal properties and displaying biodegradability have been reported.^{19–23} Nevertheless, the use of FDCA as a comonomer of succinic acid (SA) to produce fully bio-based aromatic–aliphatic copolyesters has been scarcely investigated. To our knowledge the contribution by Wu *et al.*⁶ describing a series of poly(butylene furandicarboxylate-*co*-succinate) copolyesters, the work by Jacquel *et al.*⁹ evidencing the biodegradability of some of these copolymers, and the recent paper by Lomelí-Rodríguez *et al.*²⁴ on poly(propylene furandicarboxylate-*co*-succinate) copolyesters are the only references dealing with these systems that are available in the accessible literature.

Until very recently the polymerization method invariably applied for the synthesis of both PBF and PBS and their copolyesters was the conventional melt polycondensation using either the diacid or the dimethyl ester of FDCA and SA.^{25–28} Nevertheless Loos *et al.*²⁹ have reported a series of furanic aliphatic polyesters successfully produced *via* enzymatic polycondensation of the dimethyl ester of FDCA with various potentially renewable aliphatic diols using *Candida antarctica* Lipase B (CALB).

The application of the ROP method to the synthesis of PBS was reported for the first time by Japanese researchers using as the feed a mixture of cyclic butylene succinate oligomers that were either synthesized by enzymatic cyclization^{30–32} or recovered as a subproduct in the preparation of PBS by melt polycondensation.³³ On the other hand, the first papers reporting the synthesis of PBF homopolyesters by ROP were published at the end of last year,^{34,35} and the first work using ROP for the preparation of PBF copolyesters has just been published; in that work a series of poly(butylene furandicarboxylate-*co*-terephthalate) copolyesters was synthesized and their properties evaluated as a function of the comonomer composition.³⁶

Entropically driven ring opening polymerization (ED-ROP) is a well-established technique that has been successfully used for the polymerization of cyclic oligoesters.^{37–39} The specific advantages of ROP are a high polymerization rate that allows the application of relatively mild reaction conditions and the low generation of reaction subproducts with subsequent minimization of undesirable discoloration. In this paper we applied for the first time the ED-ROP methodology to the synthesis of fully bio-based aromatic–aliphatic poly(butylene furandicarboxylate-*co*-succinate) copolyesters (*co*PBF_{*x*}S_{*y*}). The synthesis of the cyclic oligomers required for feeding has been accomplished by cyclization condensation, and their copolymerization was performed in the bulk using either tin dioctanoate (chemical process) or supported *Candida antarctica* (enzymatic process). The two ROP processes are critically compared and the two respectively produced copolyester series are extensively characterized; in particular their thermal properties are evaluated and collated to each other as well as to those reported for the same polymers obtained by conventional melt polycondensation.

2. Experimental

2.1. Materials

2,5-Furandicarboxylic acid (FDCA, >98% purity) was purchased from Satachem (China). 1,4-Butanediol (BD), thionyl chloride (SOCl₂, 99%), lipase porcine pancreas, di-azabicyclo[2.2.2]octane (DABCO, 99%) and tin dioctanoate (Sn(Oct)₂, 99%) were purchased from Sigma-Aldrich Co. Dimethyl succinate (DMS, 99%) and triethylamine (Et₃N, 98%) were purchased from Panreac. Novozym 435 (lipase B *Candida antarctica* CALB) was a gift of Novozymes. Solvents used for reaction, isolation and purification were of high-purity grade and used as received except dichloromethane (DCM), tetrahydrofuran (THF), diphenyl ether and dichlorobenzene (DCB), which were dried on 3 Å molecular sieves. DABCO catalyst was purified by sublimation.

2.2. Measurements

Elemental analyses were carried out at the Servei de Microanàlisi at IQAC (Barcelona). Tests were made in a Flash 1112 elemental microanalyser (A5), which was calibrated with appropriate standards of known composition. C and H contents were determined by the dynamic flash combustion method using He as the carrier gas. Results were given in (w/w) percentages and in duplicates. ¹H and ¹³C NMR spectra were recorded on a Bruker AMX-300 spectrometer at 25 °C, operating at 300.1 and 75.5 MHz, respectively. For NMR analysis, monomers and intermediate compounds were dissolved in deuterated chloroform (CDCl₃) and polymers in a 1 : 8 (v/v) mixture of trifluoroacetic acid (TFA) and CDCl₃, and spectra were internally referenced to tetramethylsilane (TMS). About 10 and 50 mg of the sample in 1 mL of solvent were used for ¹H and ¹³C NMR, respectively. Sixty-four scans were recorded for ¹H and between 1000 and 10 000 scans were recorded for ¹³C NMR. High-performance liquid chromatography (HPLC) analysis was performed at 25 °C in a Waters apparatus equipped with a UV detector of Applied Biosystems, operating at a wavelength of 254 nm, and a Scharlau Science column (Si60, 5 μm; 250 × 4.6 mm). Cyclic oligomers (1 mg) were dissolved in chloroform (1 mL) and eluted with hexane : 1,4-dioxane 70 : 30 (v/v) at a flow rate of 1.0 mL min⁻¹. Molecular weight analysis was performed by GPC on a Waters equipment provided with RI and UV detectors. 100 μL of 0.1% (w/v) sample solution was injected and chromatographed with a flow of 0.5 mL min⁻¹ of 1,1,1,3,3,3-hexafluoroisopropanol (HFIP). HR5E and HR2 Waters linear Styragel columns (7.8 mm × 300 mm, pore size 10³–10⁴ Å) packed with cross-linked polystyrene and protected with a precolumn were used. Molar mass average and distributions were calculated against PMMA standards. Intrinsic viscosities were measured from polymer solutions in DCA using an Ubbelohde viscometer thermostated at 25 ± 0.1 °C. Matrix-assisted laser desorption/ionization time of flight (MALDI-TOF) mass spectra were recorded in a 4700 Proteomics Analyzer instrument (Applied Biosystems) of the Proteomics Platform of Barcelona Science Park, University of Barcelona. Spectrum acquisition was performed in the MS reflector positive-ion mode. About 0.1 mg of



the sample was dissolved in 50 μL of DCM, and 2 μL of this solution was mixed with an equal volume of an anthracene solution in DCM (10 mg mL^{-1}), and the mixture was left to evaporate to dryness on the stainless steel plate of the analyser. The residue was then covered with 2 μL of a solution of 2,5-dihydroxybenzoic acid in acetonitrile:H₂O (1:1) containing 0.1% TFA, and the mixture was left to dry prior to exposure to the laser beam.

The thermal behaviour of cyclic compounds and polymers were examined by differential scanning calorimetry (DSC), using a Perkin-Elmer Pyris apparatus. The thermograms were obtained from 4–6 mg samples at heating and cooling rates of 10 $^{\circ}\text{C min}^{-1}$ under a nitrogen flow of 20 mL min^{-1} . Indium and zinc were used as standards for temperature and enthalpy calibration. The glass transition temperature (T_g) was taken as the inflection point of the heating DSC traces recorded at 20 $^{\circ}\text{C min}^{-1}$ from melt-quenched samples, and the melting temperature (T_m) was taken as the maximum of the endothermic peak appearing on heating traces. Thermogravimetric analysis was performed on a Mettler-Toledo TGA/DSC 1 Star System under a nitrogen flow of 20 mL min^{-1} at a heating rate of 10 $^{\circ}\text{C min}^{-1}$ and within a temperature range of 30 to 600 $^{\circ}\text{C}$.

X-Ray diffraction patterns from powdered samples coming directly from synthesis were recorded on a PANalytical X'Pert PRO MPD θ/θ diffractometer using $\text{CuK}\alpha$ radiation of wavelength 0.1542 nm. Micrographs were recorded using an Olympus BX 51 polarizing optical microscope provided with a Linkam THMS 600 stage. For observation, 10 mg of each sample were dissolved in 1 mL of chloroform or HFIP, and aliquots of 0.2 mL were slowly evaporated on a glass slide.

2.3. Synthesis of cyclic oligomers

$c(\text{BF})_n$ oligomers. Cyclic butylene 2,5-furandicarboxylate oligomers $c(\text{BF})_n$ were synthesized by high-dilution condensation (HDC) from mixtures of BD and 2,5-furandicarboxylic dichloride (FDCA-Cl₂) as previously reported.³⁵ Briefly, to a three-necked round-bottom flask charged with 250 mL of THF and cooled to 0 $^{\circ}\text{C}$, 12.5 mmol (1.40 g) of DABCO was added under stirring. 5 mmol (0.96 g) of FDCA-Cl₂ and 5 mmol (0.46 g) of BD in 10 mL of THF were added drop-wise simultaneously for 40 min using separated addition funnels in order to maintain the equimolarity of the reagents in the reaction mixture. The reaction was finished by adding 1 mL of water followed by 5 mL of 1 M HCl, and after stirring for 5 min, the reaction mixture was diluted with DCM and filtered. The filtrate was washed with 0.1 M HCl, dried on MgSO_4 , and evaporated to dryness to render a mixture of linear and cyclic oligomers. Linear oligomers were removed by chromatography through a short column of silica gel using a cold mixture of DCM/diethyl ether 90:10 (v/v) as the eluent. Yield 75%. ¹H NMR (δ ppm, CDCl_3 , 300 MHz): 7.25 (s, 2H), 4.40 (t, 4H), 1.92 (q, 4H). ¹³C NMR (δ ppm, CDCl_3 , 75.5 MHz): 157.9, 146.5, 118.7, 64.8, 25.4.

$c(\text{BS})_n$ oligomers. Cyclic butylene succinate oligomers $c(\text{BS})_n$ were obtained by enzymatic cyclization of BD and DMS following the procedure reported by Sugihara *et al.*³⁰ with slight

modifications. Briefly, a three-necked round-bottom flask was charged with 250 mL of toluene, 4.97 mmol (0.73 g) of DMS, 5.07 mmol (0.46 g) of BD and 1.184 g of CALB (100% w/w of the relative total monomer concentration) and the mixture was left to react for 48 h at 96 $^{\circ}\text{C}$ under a nitrogen flow. 4 Å molecular sieves were placed at the top of the flask to absorb the released methanol. The reaction mixture was then diluted with 50 mL of CHCl_3 and the enzyme was removed by filtration. The solvent was evaporated under reduced pressure to obtain the oligomer mixture, which was analysed by NMR and MALDI-TOF. Yield: 80%. ¹H NMR (δ ppm, CDCl_3 , 300 MHz): 4.14 (m, 4H), 2.64 (m, 4H), 1.72 (m, 4H). ¹³C NMR (δ ppm, CDCl_3 , 75.5 MHz): 171.90, 64.27, 29.37, 26.13.

2.4. Synthesis of polymers

Chemically catalyzed ROP in the bulk. Mixtures of $c(\text{BF})_n$ and $c(\text{BS})_n$ at different molar ratios were polymerized following the same procedure we previously used for the synthesis of PBF and P(BF-co-BT) copolyesters.³⁶ A total of 47 mmol of the mixture of cyclic oligomers with the selected composition together with 0.5 mol% of $\text{Sn}(\text{Oct})_2$ were dissolved in 10 mL of DCM and dried under vacuum at room temperature for 24 h. The mixture was then placed in a three-necked round-bottom flask provided with mechanical stirring and left to react at five different temperatures (180, 200, 210, 220 and 230 $^{\circ}\text{C}$) for 12 h. Aliquots were withdrawn at scheduled times and analysed by GPC. The NMR spectra of the ensuing copolyesters are available in the ESI (Fig. SI-1†).

Enzymatically catalyzed ROP in the bulk. Ring opening polymerization of the $c(\text{BF})_n$ and $c(\text{BS})_n$ mixtures was performed in the bulk with the assistance of CALB. Briefly, 47 mmol of the cyclic oligomer mixture was placed in a three-necked round-bottom flask provided with a nitrogen flow and heated at five different temperatures (120, 130, 140, 145 and 150 $^{\circ}\text{C}$) under stirring. Then CALB (40% w/w) was added to the molten mass and the reaction was left to proceed for 24 h. The reaction mixture was then cooled down to room temperature and a large volume of CHCl_3 was added to dissolve the polymer. After removing the enzyme by filtration, the solvent was evaporated under vacuum and the solid residue was dried for 24 h before analysis. Aliquots were withdrawn at scheduled times and analysed by GPC. The NMR spectra of the copolyesters thus obtained are available in the ESI (Fig. SI-2†).

Blank assays were carried out for comparative purposes. For this, a mixture of 0.24 mmol (49.3 mg) of $c(\text{BF})_n$ and 0.94 mmol (161.7 mg) of $c(\text{BS})_n$ without any catalyst added was made to react at 150 and 220 $^{\circ}\text{C}$ for 12 and 24 h, respectively. The evolution of the reaction was followed as above and the resulting products were analysed by NMR.

Enzymatically catalyzed ROP in solution. For comparative purposes, the copolyester $\text{coPBF}_{50}\text{S}_{50}$ was also synthesised by ROP in solution assisted by CALB. Briefly, a mixture of 0.29 mmol (60.9 mg) of $c(\text{BF})_n$ and 0.29 mmol (50 mg) of $c(\text{BS})_n$ was dispersed in diphenyl ether (1 g) and the mixture was placed in a three-necked round-bottom flask under stirring. When the mixture was solubilized, 40% w/w of CALB was



added. The reaction was conducted at 50 °C for 24 h and aliquots were withdrawn at scheduled times to follow the advance of the reaction by GPC analysis. Finally the reaction mass was diluted by adding CHCl₃, and CALB was removed by filtration. The clean solution was concentrated and precipitated with hexane. The recovered copolyester was dried in vacuum for 2 days before being subjected to analysis.

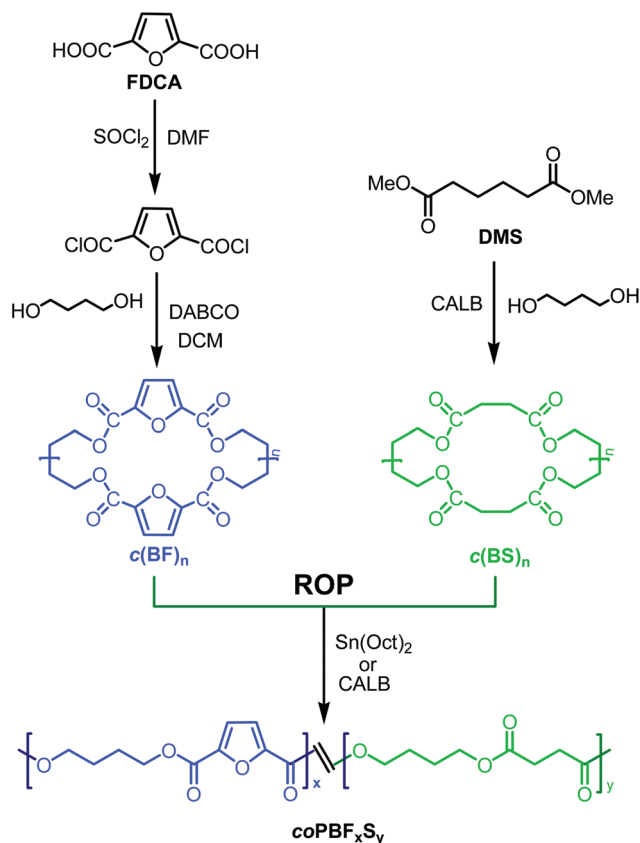
Hydrolytic degradation and biodegradation. Films for hydrolytic degradation and biodegradation studies were prepared from PBF, *co*PBF₆₀S₄₀, *co*PBF₄₀S₆₀ and PBS with a thickness of ~200 μm by casting from either HFIP or chloroform solution at a polymer concentration of 100 g L⁻¹. The films were cut into 10 mm-diameter, 20–30 mg-weight disks and dried under vacuum to a constant weight. For hydrolytic degradation, samples were immersed in vials containing 10 mL of either citric acid buffer, pH 2.0 or sodium phosphate buffer, pH 7.4 at 37 °C. Enzymatic degradation assays were carried out at 37 °C in pH 7.4 buffered sodium phosphate solution with 10 mg of lipase porcine pancreas added. In this case the buffered solution was replaced every 72 h in order to maintain the enzyme activity. In both cases, the disks were withdrawn from the incubation medium at scheduled periods of time, washed carefully with distilled water, dried to a constant weight and finally analysed by GPC.

3. Results and discussion

The strategy followed in this work to produce the poly(butylene 2,5-furandicarboxylate-*co*-succinate) copolyesters is depicted in Scheme 1. Firstly cyclic oligomers of both butylene 2,5-furandicarboxylate, *c*(BF)_{*n*}, and butylene succinate, *c*(BS)_{*n*}, were synthesized, and secondly, mixtures of the two types of cycles were polymerized through ring opening reactions catalysed either chemically or enzymatically.

3.1. Synthesis of cyclic oligomers

In a recent study we reported for the first time the synthesis of cyclic (butylene 2,5-furandicarboxylate) oligomers by both high dilution condensation (HDC) and cyclodepolymerization (CD) methods. Both methods were suitable for *c*(BF)_{*n*} synthesis but the former afforded the oligomeric mixture in slightly higher yield and more enriched with the cyclic dimer.³⁵ In the present work we have made use of the HDC method to produce *c*(BF)_{*n*}, with the results shown in Table 1. After purification by column chromatography, the ¹H NMR analysis (Fig. 1a) was in full agreement with the expected structure and showed no signal arising from linear species. The HPLC chromatogram (Fig. 1b) displayed three peaks corresponding to cyclic dimers, trimers and tetramers in an approximate 8 : 4 : 1 ratio, although the MALDI-TOF analysis (Fig. 1c) revealed the presence of cyclic species of higher sizes in minor amounts. Nevertheless, this mixture without further fractionation was used for ROP since our previous kinetics study on the polymerization of *c*(BF)_{*n*} showed no relevant differences when cycles of different sizes or their mixtures were used for feeding.³⁵



Scheme 1 Synthesis route to poly(butylene furandicarboxylate-*co*-succinate) copolyesters by ROP.

Table 1 Cyclization reaction results

	Method ^a	Yield (%)	Composition ^b (2/3/4)	<i>T</i> _m ^c (°C)	^o <i>T</i> _{5%} ^c (°C)	<i>T</i> _d ^c (°C)
<i>c</i> (BF) _{<i>n</i>}	HDC	67	61/31/8	147	276	387
<i>c</i> (BS) _{<i>n</i>}	EC	70	50/40/10	105	266	340

^a HDC: high dilution cyclization using DABCO/Et₃N; EC: enzymatic cyclization using CALB. ^b Relative content (w/w) of the reaction product in cyclic dimers, trimers and tetramers as measured by HPLC. ^c Melting and decomposition temperatures measured by DSC and TGA.

The synthesis of *c*(BS)_{*n*} was enzymatically performed as well, and the obtained results are given in Table 1. We applied the method reported by Sugihara *et al.*³⁰ which consists of cyclizing a mixture of BD and DMS dissolved in toluene by the action of CALB. The ¹H NMR and MALDI-TOF spectra as well as the HPLC chromatogram of the cyclization reaction product without being subjected to purification are shown in Fig. 2. The mixture is composed almost exclusively of cyclic species with sizes ranging from dimers to nonamers; cyclic dimers, trimers and tetramers are in an approximate 5 : 4 : 1 ratio, and their sum amounts to more than 90% of the total oligomers present in the mixture. Here again the oligomer mixture without fractionation was used for ROP since



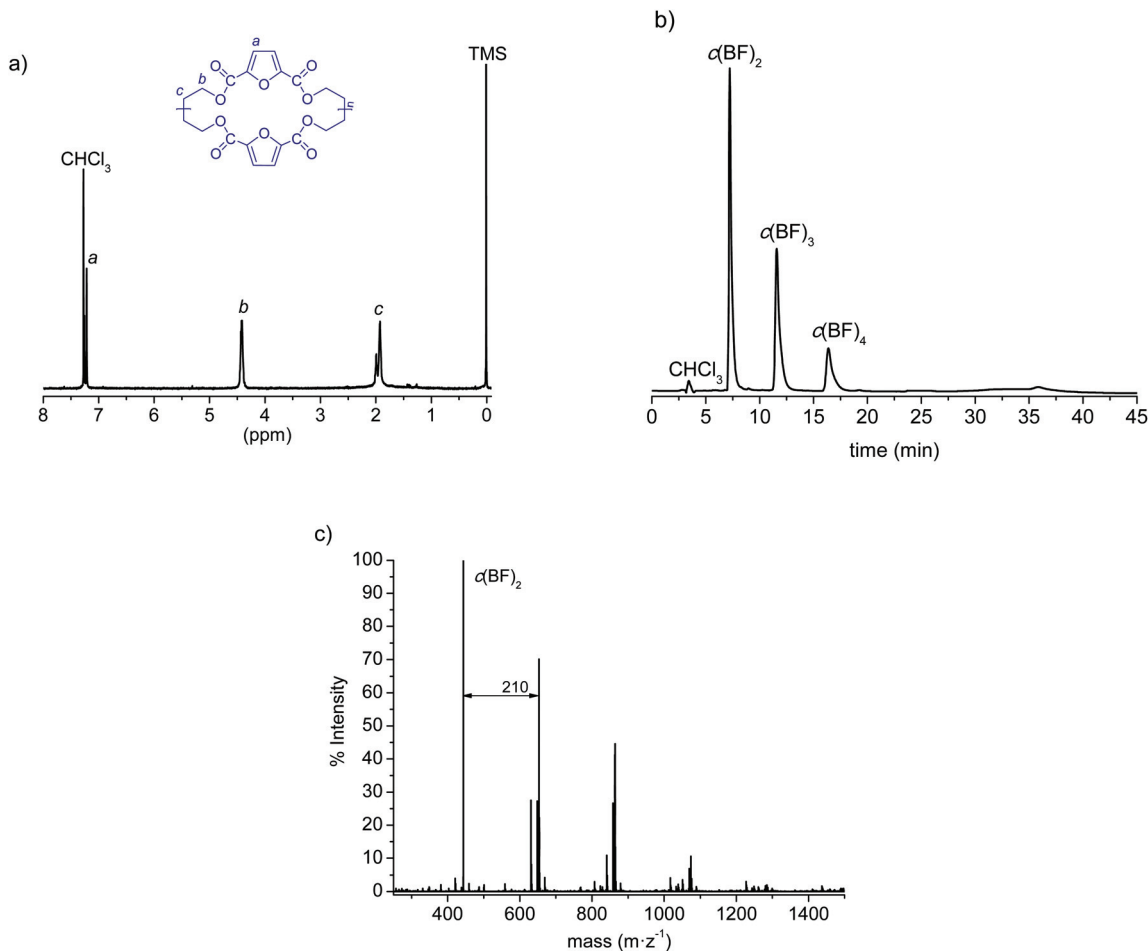


Fig. 1 (a) ¹H NMR, (b) HPLC and (c) MALDI-TOF spectra for c(BF)_n.

kinetics studies carried out by Kondo *et al.*⁴⁰ have shown that polymerization results are not substantially affected by the cycle size.

Both c(BF)_n and c(BS)_n mixtures are white semicrystalline powders with melting temperatures of 147 °C and 105 °C, respectively. They start to decompose by the effect of heating at temperatures above 250 °C, following a single-step degradation process that displays its maximum rate at temperatures between 300 and 400 °C. The TGA traces recorded from these mixtures along with their derivative curves are available in the ESI (Fig. SI-3†).

3.2. Synthesis of copBF_xS_y copolyesters

3.2.1. ROP catalysed by tin octanoate. The polymerization of mixtures of c(BF)_n and c(BS)_n catalysed by tin octanoate was performed at temperatures above the melting of the oligomer mixtures. The effect of temperature on the polymerization rate was indirectly evaluated by measuring the molecular weight of the polyesters obtained after 12 h of reaction at different temperatures. Three c(BF)_n/c(BS)_n ratios (80 : 20, 50 : 50 and 20 : 80) in addition to the two pristine cyclic oligomer fractions were examined. The results are shown in Fig. 3a, where *M*_w is

plotted against six temperatures increasing 20 or 10 °C from 180 up to 230 °C. In general a steady increase in the molecular weight of the resulting polymer was observed for all compositions as the temperature was raised. It was noticeable however that the *M*_w of both the copolyester copBF₈₀S₂₀ and the homopolyester PBF levelled off at temperatures above ~200 °C, probably due to the higher thermal sensitivity to degradation of the furan-based compounds. Nevertheless, the relevant outcome of this study is that the molecular weight of the resulting polyester increases with the content of the feed in c(BF)_n at whichever reaction temperature. This result brings into evidence the higher reactivity of the furan-based cyclic oligomers in chemically catalysed ROP.

The evolution of the molecular weight along the reaction time was then examined for the polymerization of the same oligomer cyclic mixtures carried out at 220 °C, and the obtained results are shown in Fig. 3b. The trends shown by PBF and PBS are remarkably different; the c(BF)_n fraction polymerized initially very quickly to reach an almost constant molecular weight of about 65 000 g mol⁻¹ after 2–3 h of reaction, whereas c(BS)_n polymerized following a nearly linear trend to render a PBS with a *M*_w near 45 000 g mol⁻¹ after 12 h



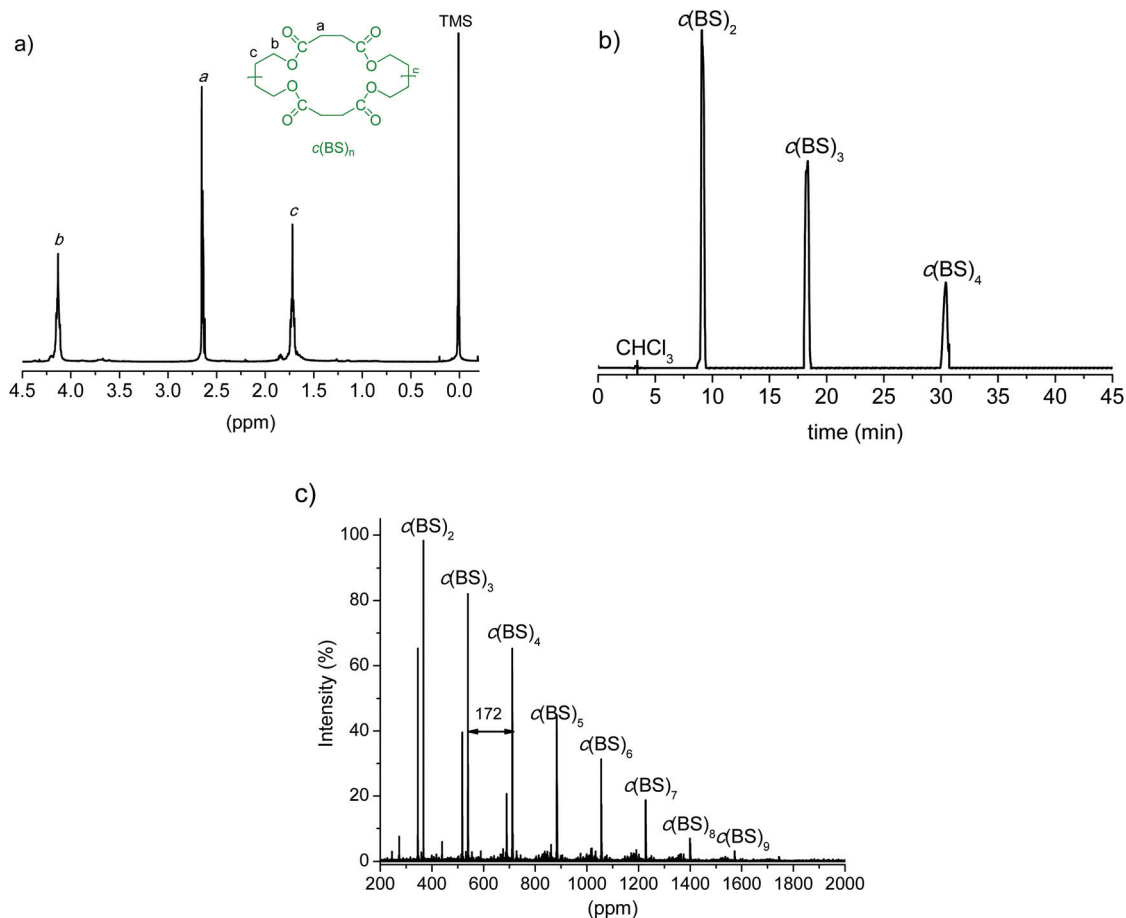


Fig. 2 (a) ^1H NMR, (b) HPLC and (c) MALDI-TOF spectra for $c(\text{BS})_n$.

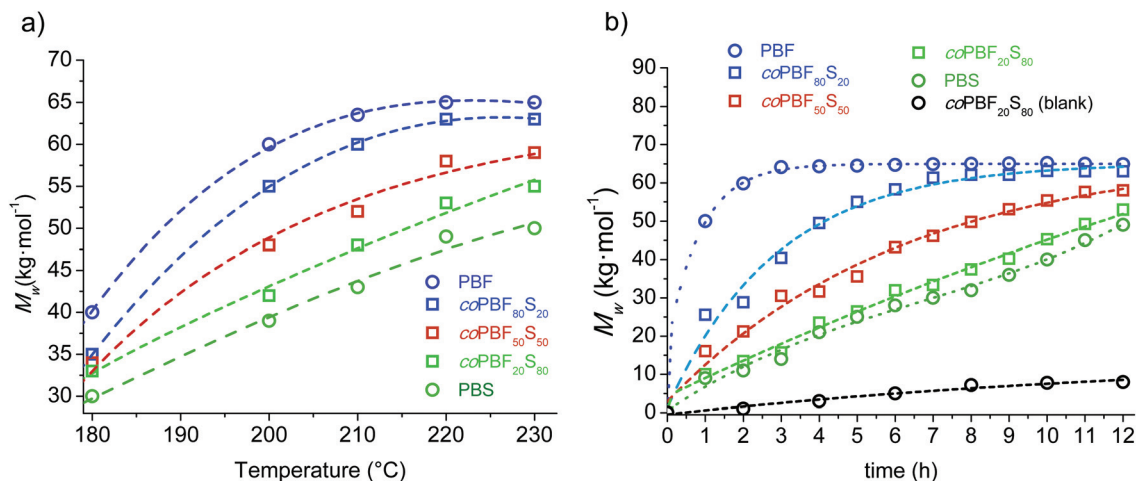


Fig. 3 Effect of temperature and time on $\text{Sn}(\text{Oct})_2$ catalyzed ROP of mixtures of $c(\text{BF})_n$ and $c(\text{BS})_n$ carried out in the melt. (a) M_n of coPBF_xS_y against reaction temperature after 12 h of reaction time. (b) Evolution of M_n with reaction time at 220 °C.

of reaction. The behaviour observed for the copolyesters were intermediate between those displayed by the two homopolymers, with the profile being closer to one or the other according to the composition.

3.2.1. ROP catalysed by *Candida antarctica*. The ROP of mixtures of $c(\text{BF})_n$ and $c(\text{BS})_n$ catalysed by lipase *Candida antarctica* was carried out in the bulk with the fixed CALB dispersed in the molten reaction mass. The kinetics study carried



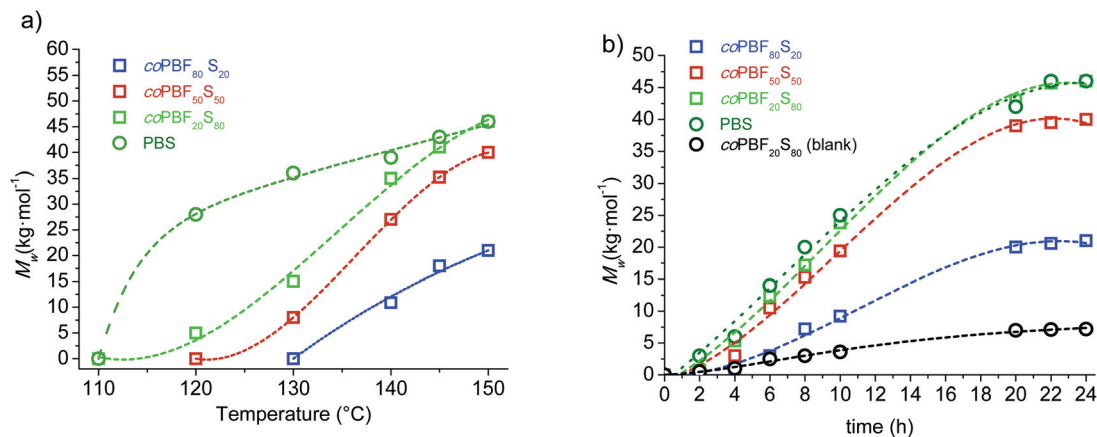


Fig. 4 Effect of temperature and time on CALB assisted ROP of mixtures of $c(\text{BF})_n$ and $c(\text{BS})_n$ in the melt. (a) M_w of PBF_xS_y against reaction temperature for a reaction time of 24 h. (b) Evolution of M_w with reaction time at 150 $^{\circ}\text{C}$.

out on this system was similar to that described above for the chemically catalysed ROP, *i.e.* the influence of both temperature and time on the molecular weight of the growing polyester was evaluated for different compositions. However, the assay conditions had to be adapted to the limitations imposed by the use of the enzyme. The temperature had to be restricted to the 120–150 $^{\circ}\text{C}$ range because below 120 $^{\circ}\text{C}$ the mixture was not fully molten and above 150 $^{\circ}\text{C}$ the enzyme activity was rapidly lost. This prevented the preparation by this method of polyesters with melting temperatures above 150 $^{\circ}\text{C}$ as is the case for both PBF and the copolyester $coPBF_{90}S_{10}$. The results obtained after 24 h of reaction at temperatures of 120, 130, 140, 145 and 150 $^{\circ}\text{C}$ are plotted in Fig. 4a.

The trend followed by the molecular weight with increasing reaction temperature was not so predictable as in the case of chemocatalyzed ROP. The profile shape changed significantly with the composition, and differences among the M_w values largely vary with temperature. Notwithstanding, it was interestingly noticed that polymer growing catalyzed by *C. antarctica* was more favoured as the content of butylene succinate increased, which is exactly opposite to the behaviour observed in tin catalysed reactions. In fact, minimum M_w values were reached at all assayed temperatures for the $\text{PBF}_{80}\text{S}_{20}$, which is the copolyester with the highest content of butylene furandicarboxylate units attainable by this method.

The evolution of the molecular weight of the polyesters with time in the CALB assisted ROP of $c(\text{BF})_n/c(\text{BS})_n$ mixtures at 150 $^{\circ}\text{C}$ along a period of 24 h is depicted in Fig. 4b. The M_w vs. time profiles are of similar shape for the four assayed polyesters with the slope decreasing with the content of BF units. Differences in the polymerization rates become more apparent as the reaction progresses, and the molecular weights reached at the end of the process were lower or higher than those obtained by chemocatalysis depending on the composition.

In order to appraise the possible occurrence of self-polymerization of the $c(\text{BF})_n/c(\text{BS})_n$ by ring opening reaction

induced by heating, a series of blank assays, *i.e.*, assays conducted under the same reaction conditions but in the absence of any catalyst, was carried out with the mixture $c(\text{BF})_n/c(\text{BS})_n$ (20 : 80). The profiles M_w vs. either temperature or time that were obtained from these assays under chemically or enzymatically catalyzed conditions are included in Fig. 3b and 4b for comparison. The highest molecular weight reached in these assays was about 5000 $\text{g}\cdot\text{mol}^{-1}$, corresponding to oligomers of about 10 monomeric units. The ^1H NMR spectra of the feeding mixture and the reaction product of the ROP carried out either without catalyst or in the presence of CALB are compared in Fig. 5. This comparison reveals that $c(\text{BF})_n$ remained unreacted in the blank whereas an appreciable amount of $c(\text{BS})_n$ cycles disappeared. Conversely, both $c(\text{BF})_n$ and $c(\text{BS})_n$ reacted almost completely in the presence of CALB.

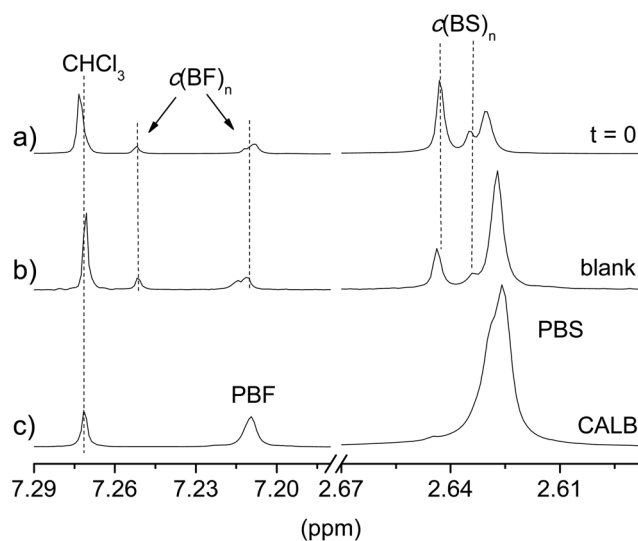


Fig. 5 Enlarged 7.3–7.18 and 2.68–2.60 ppm regions of the ^1H NMR spectra of the initial $c(\text{BF})_n/c(\text{BS})_n$ (20 : 80) mixture (a), after reaction at 150 $^{\circ}\text{C}$ for 24 h in the absence of catalyst (b), and with CALB added (c).



3.3. Chemical structure and microstructure of copolyesters

The chemical structure, including both composition and microstructure as well as the molecular weight, of the homopolyesters and copolyesters obtained by the two ROP methods applied in this work were systematically determined, and results are gathered in Table 2. These data refer to polyesters synthesized with Sn(Oct)₂ as the catalyst at 230 °C for 12 h, and to those synthesized with the assistance of CALB at 150 °C for 24 h. The chemical constitution of all the polyesters was ascertained by NMR spectroscopy. The ¹H NMR spectra of the copolyester PBF₅₀S₅₀ prepared by either chemo-ROP or enzym-ROP are shown in Fig. 6 for illustrative purposes. Both spectra are almost undistinguishable with all peaks assigned and displaying no signals other than those expected for the predicted chemical constitution. Both the ¹H and ¹³C NMR spectra of the two whole sets of synthesized polyesters are given in the ESI (Fig. SI-1 and SI-2†).

The molar composition in BF and BS units of the two series of copolyesters was determined by quantification of the signals *b* and *b'* arising from the α-methylene of the butylene furandicarboxylate and butylene succinate units, respectively, and the obtained values are listed in Table 2. In support of the NMR determination, the elemental analysis of the copolyester *co*PBF₆₀S₄₀ obtained by the two methods was carried out as an example. The results were fully consistent with the predicted chemical constitution and composition determined by NMR for both cases (Table 2).

A comparative inspection of the feed and copolyester compositions reveals that deviations between them are higher

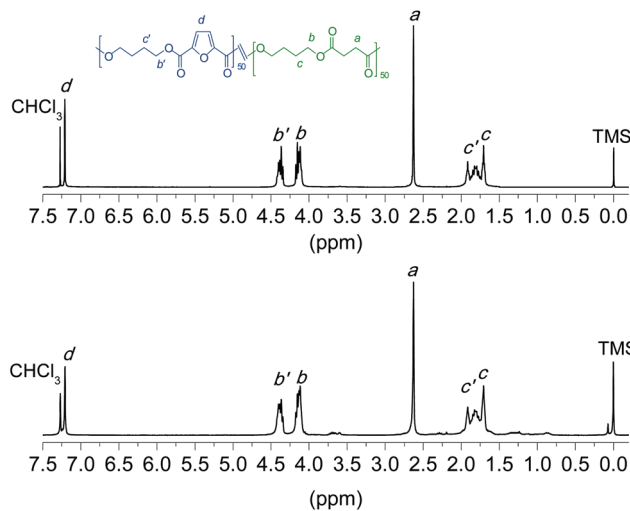


Fig. 6 ¹H NMR spectra of *co*PBF₅₀S₅₀ obtained by chemically (top) and enzymatically (bottom) catalyzed ROP.

when the ROP is chemically catalysed, and that in this case, they correspond to BS unit losses. Furthermore, the differences become more noticeable as the content of BF increases, with the loss in BS becoming around 40% for the *co*PBF₉₀S₁₀ copolyester. Conversely, the observed deviations were less than 5% for the copolyesters synthesized by enzymatically assisted ROP. These results are consistent with the different relative reactivity of the *c*(BF)_{*n*} and *c*(BS)_{*n*} species in the two polymerization methods, although reasons due to differences in

Table 2 Yields, compositions, average molecular weights, and microstructure of *co*PBF_{*x*}S_{*y*} copolyesters obtained via chemical and enzymatic ROP^a

Copolyester	Yield (%)	Composition ^b X _{BF} /Y _{BS} (mol mol ⁻¹)	Molecular weight ^c			Dyad content ^d (mol%)			Average sequence length ^e		R ^e
			M _w (g mol ⁻¹)	D	[η] (dL g ⁻¹)	FBF	FBS + SBF	SBS	n _{BF}	n _{BS}	
PBF	95	100/0	65 000	1.98	0.76	—	—	—	—	—	—
	—	100/0	—	—	—	—	—	—	—	—	—
<i>co</i> PBF ₉₀ S ₁₀	92	94/6	64 000	1.79	0.74	93.2	2.6	4.2	6.58	1.30	0.92
	—	—	—	—	—	—	—	—	—	—	—
<i>co</i> PBF ₈₀ S ₂₀	93	84/16	63 000	1.85	0.74	66.7	28.2	5.1	5.73	1.36	0.91
	65	81/19	21 000	1.30	0.25	67.8	25.2	7.0	6.38	1.56	0.80
<i>co</i> PBF ₆₀ S ₄₀	90	65/35 ^f	62 000	1.90	0.71	39.5	48.8	11.7	2.62	1.48	1.06
	70	58/42 ^f	23 000	1.30	0.30	35.3	45.1	19.6	2.57	1.87	0.92
<i>co</i> PBF ₅₀ S ₅₀	90	51/49	58 000	1.79	0.62	27.0	48.6	48.7	2.11	2.00	0.97
	91	49/51	40 000	1.60	0.55	15.7	47.6	36.6	1.66	2.53	0.99
<i>co</i> PBF ₄₀ S ₆₀	90	43/57	56 000	1.95	0.65	16.7	52.2	31.1	1.64	2.19	1.07
	92	41/59	43 000	1.70	0.55	13.4	41.3	65.9	1.65	3.19	0.91
<i>co</i> PBF ₂₀ S ₈₀	89	22/78	53 000	1.90	0.54	5.2	31.3	63.5	1.33	5.06	0.95
	91	22/78	46 000	1.50	0.53	3.2	33.7	63.0	1.19	4.73	1.05
<i>co</i> PBF ₁₀ S ₉₀	88	12/88	53 000	1.89	0.60	0	22.6	77.4	1.00	7.8	1.13
	91	11/89	47 000	1.50	0.53	1.3	15.6	83.1	1.17	11.6	0.94
PBS	95	0/100	49 000	1.97	0.55	—	—	—	—	—	—
	92	0/100	46 000	1.40	0.51	—	—	—	—	—	—

^a Copolyesters synthesized using Sn(Oct)₂ (top row) and CALB (bottom row). ^b Copolyester compositions determined from signals observed in the ¹H NMR spectra. ^c M_w, D, and η determined by GPC and Ubbelohde viscosity, respectively. ^d Determined by integration of the α-methylene signals in the butylene segments in the ¹³C NMR spectra. ^e Average lengths of homogeneous BF and BS sequences and randomness (R) calculated by applying a statistical approach.⁴¹ ^f Calculated for *co*PBF₆₅S₃₅: C, 56.73%; H, 5.47%; Found: C, 53.37%; H, 5.54%. Calculated for *co*PBF₅₈S₄₂: C, 56.40%; H, 5.67%; Found: C, 56.06%; H, 5.96%.



volatility should not be fully neglected provided that much higher temperatures are used in the chemical procedure.

The molecular weights measured by GPC against polystyrene standards were in the 65 000–50 000 and 47 000–20 000 g mol⁻¹ ranges for polyesters obtained by chemically and enzymatically catalysed ROP, with dispersities significantly lower in the latter case. Intrinsic viscosities were in agreement with the M_w values determined by GPC. The trends observed for the molecular weights in the two series are opposite to each other according again to the different relative reactivities displayed by $c(\text{BF})_n$ and $c(\text{BS})_n$ in each of the two polymerization methods.

The microstructure of the copolyesters was elucidated by ¹³C NMR analysis, taking benefit of the sensitivity of the α -carbon of the butylene segment to the distribution sequence at the dyad level. The four possible dyads, FBF, SBS, and FBS/SBF, appear within the 60.5–58.5 ppm region (Fig. SI-4†) as four well-resolved peaks whose areas were integrated for each copolyester composition, and the resulting values used in the corresponding equations to calculate the number average sequence length. The results are listed in Table 2 for the two copolyester series, where it can be seen that the n_{BF} and n_{BS} values vary steadily with the composition in opposite directions, and both approximate to 2 for the 50 : 50 composition. The degree of randomness is between 0.9 and 1.1, indicating that an approximate random distribution of BF and BS units is present in all the copolyesters. The conclusion that a coPBF_xS_y random copolyester is genuinely generated in the ROP of mixtures of $c(\text{BF})_n$ and $c(\text{BS})_n$ whether it is chemically or enzymatically catalyzed cannot be straightforwardly drawn from such results since randomization by transesterification is very probable to occur at the high temperatures used in these processes. In order to clarify this point, ROP of the $c(\text{BF})_n/c(\text{BS})_n$ (50 : 50) mixture assisted by CALB was accomplished in solution at 50 °C, a temperature at which the occurrence of transesterification reactions may be neglected. The $\text{coPBF}_{50}\text{S}_{50}$ copolyester resulting in these conditions was obtained in a 80% yield and had a M_w of 14 000 g mol⁻¹ and a D of 1.8. Its ¹³C NMR spectrum was similar to that recorded from the copolyester obtained a high temperature; specifically, no differences can be appreciated when they are compared in the 60.5–58.5 ppm region (Fig. 7, inset). It can be therefore concluded that the ROP of $c(\text{BF})_n/c(\text{BS})_n$ renders random copolyesters whatever the applied reaction conditions.

3.4. Thermal properties

The thermal properties of the polyesters and copolyesters composing the two series prepared by ROP were comparatively examined by TGA and DSC, and the most relevant parameters resulting from the study are collected in Table 3. Moreover, data are also compared with those reported by Wu *et al.*, which were obtained from polyesters with similar compositions but prepared by two-step melt polycondensation.⁶ The TGA weight loss– T traces of all the polyesters synthesized in this work as well as their derivative curves are available in the ESI (Fig. SI-5†). In all cases thermal

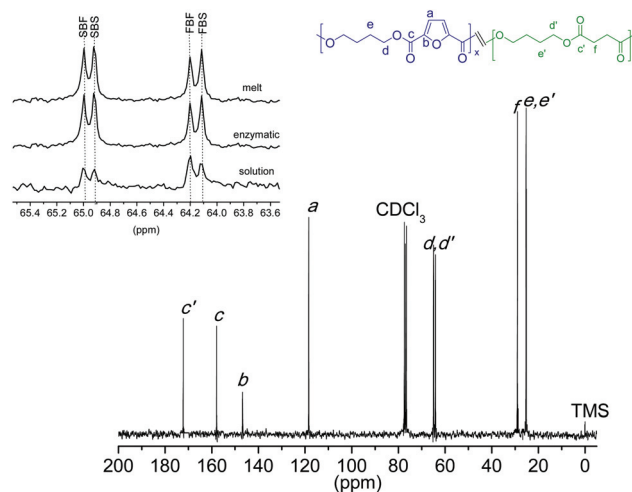


Fig. 7 ¹³C NMR spectra of $\text{coPBF}_{50}\text{S}_{50}$ obtained by ROP at 50 °C in solution catalyzed by $\text{Sn}(\text{Oct})_2$. Inset: enlarged 60.5–58.5 ppm region compared for different methods of polymerization.

decomposition proceeded through a single step taking place within the 390–400 °C range and without significant differences between the two series. The onset decomposition temperatures of the copolyesters were between 350 and 340 °C, and polymers prepared by enzymatic ROP invariably started to decompose at a few degrees below those obtained by ROP catalysed by $\text{Sn}(\text{Oct})_2$, which is reasonable due to their lower molecular weight. The residue left after heating at 600 °C under a nitrogen flow was significantly smaller in the enzymatically synthesized polyesters. This is a much-expected result because of the absence of metallic components in these polymers. The thermal decomposition data obtained for the ROP-synthesized coPBF_xS_y are essentially coincident with those reported for this family of copolyesters obtained by polycondensation, with slight differences in magnitudes being explainable by differences in molecular weight or experimental inaccuracy.

The thermal transitions, *i.e.* glass and melting/crystallization, were examined by DSC. The DSC traces of the two series registered at heating and cooling from the melt are included in the ESI (Fig. SI-6†), and the measured thermal parameters are listed in Table 3. All the pristine copolyesters displayed crystallinity with enthalpy values decreasing as the content of BF and BS units becomes more equilibrated, so that only those copolyesters containing more than 40% of one or another unit are able to crystallize from the melt. A representative selection of polarizing optical micrographs showing the crystalline textures displayed by these copolyesters upon crystallization is available in the ESI (Fig. SI-7†). The XRD analysis of the copolyesters gave results in agreement with the DSC data. The scattering profiles registered from crystalline coPBF_xS_y and from the two homopolyesters PBF and PBS prepared by chemically catalyzed ROP are compared in Fig. 8. It is clearly seen that the characteristic scattering produced by one or another homopolyester is present in the copolyester profiles, with more



Table 3 Thermal properties of coPBF_xS_y ^a

Copolymer	TGA ^b			DSC ^c				
	^o T _d (°C)	^{max} T _d (°C)	W (%)	T _g (°C)	T _m ¹ (°C)	ΔH _m ¹ (J g ⁻¹)	T _m ² (°C)	ΔH _m ² (J g ⁻¹)
PBF	370	400	7	42	170	40	170	35
<i>coPBF</i> ₉₀ <i>S</i> ₁₀	350	395	8	28	166	38	164	30
<i>coPBF</i> ₈₀ <i>S</i> ₂₀	350	395	7	23	146	33	146	27
<i>coPBF</i> ₆₀ <i>S</i> ₄₀	348	396	5	20	140	15	140	17
<i>coPBF</i> ₆₀ <i>S</i> ₄₀	343	397	6	7	114	13	—	—
<i>coPBF</i> ₅₀ <i>S</i> ₅₀	342	397	5	5	126	7	—	—
<i>coPBF</i> ₅₀ <i>S</i> ₅₀	349	400	5	-1	95	11	—	—
<i>coPBF</i> ₄₀ <i>S</i> ₆₀	341	399	1	-6	86	4	—	—
<i>coPBF</i> ₄₀ <i>S</i> ₆₀	347	399	6	-13	60	6	—	—
<i>coPBF</i> ₄₀ <i>S</i> ₆₀	344	399	2	-20	72	10	—	—
<i>coPBF</i> ₂₀ <i>S</i> ₈₀	343	399	5	-23	90	55	93	32
<i>coPBF</i> ₂₀ <i>S</i> ₈₀	338	400	3	-23	93	30	93	51
<i>coPBF</i> ₁₀ <i>S</i> ₉₀	340	401	3	-23	100	71	105	49
<i>coPBF</i> ₁₀ <i>S</i> ₉₀	337	400	2	-25	102	55	103	63
PBS	350	399	3	-30	112	106	114	82
PBS	348	398	3	-40	112	60	111	87

^a Values in the upper and lower rows correspond to polyesters obtained by tin and lipase catalyzed ROP, respectively. ^b ^oT_d is the onset degradation temperature for 5% of weight lost and ^{max}T_d is temperature at which degradation happens at the maximum rate. ^c 1 and 2 refer to data registered at the first heating and second heating, respectively.

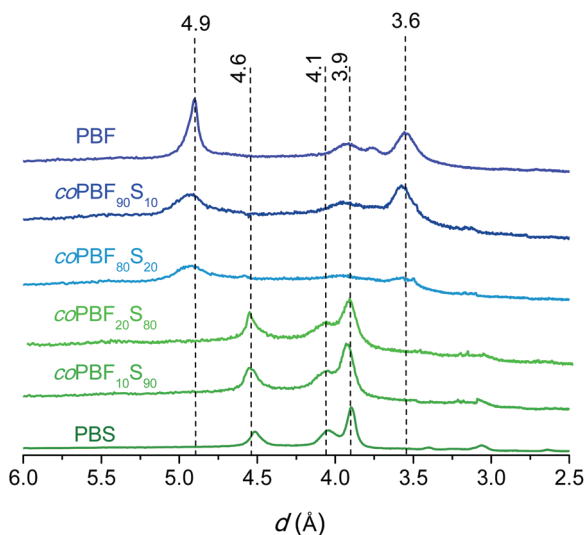


Fig. 8 Powder X-ray diffraction profiles of coPBF_xS_y synthesized by ROP with $\text{Sn}(\text{Oct})_2$.

or less intensity depending on the composition. For BF/BS ratios close to 1, the crystallinity becomes so low that no discrete scattering is perceivable on the XRD profiles of the copolymers. As it is commonly found, the crystal structure of the homopolyesters is fully retained by the copolymers containing limited amounts of the second comonomer, beyond which they become essentially amorphous.

The melting and glass-transition temperatures of the coPBF_xS_y copolymers are properties of high relevance regarding their possible applications as thermoplastics. The consequence of incorporating the furanoate units in the PBS is

to modify both the T_m and T_g . The variation of T_m with composition for the two series of coPBF_xS_y copolymers respectively prepared by the two ROP methods here described is plotted in Fig. 9a. In this plot, it is clearly shown how T_m decreases along the BS-high composition range to invert the trend when the copolymer starts to become enriched with BF units. This behavior is very similar for the two series, and is also similar to that described for the same copolymers prepared by melt polycondensation, with the slight deviations observed among them being attributable to differences in molecular weights or of experimental origin. A similar trend is observed for the melting enthalpy, which is readily explained by the incompatibility of BF and BS to cocrystallize. Nevertheless it is worthy to notice the presence of some crystallinity in the copolymers containing approximately similar amounts of the two units, which brings into evidence the ability of BF and BS short sequences for crystallizing. Similar behavior has been recently reported for random copolymers made of butylene furanoate and butylene terephthalate units.³⁶

All the copolymers display a single T_g in agreement with the random monomer distribution determined by NMR. The variation of T_g with the composition is plotted in Fig. 9b for the two series as well as for the series made by melt polycondensation reported by Wu *et al.*⁶ In all cases the T_g of PBS goes up continuously with the content of BF units, which should be expected for the increase in chain stiffness. These results demonstrate the possibility of tuning the T_g of the copolymers by adjusting the composition. Again small discrepancies are observed among copolymers produced by different procedures. Such deviations become more apparent for the copolymers with intermediate compositions produced



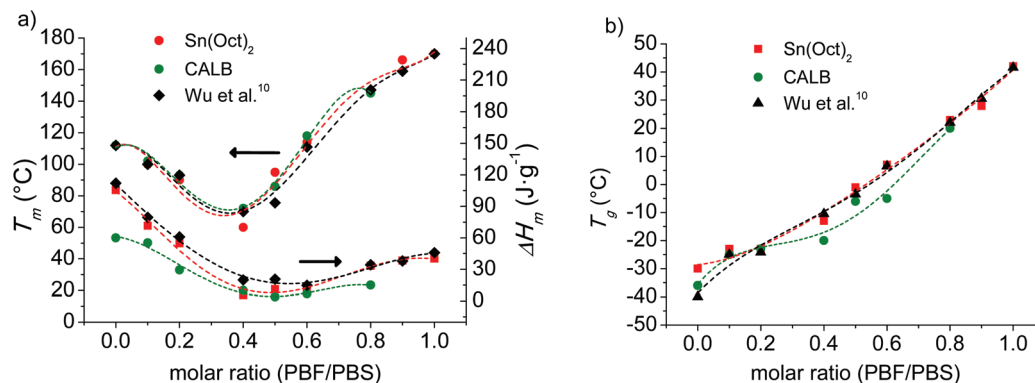


Fig. 9 T_m (a) and T_g (b) of coPBF_xS_y as a function of composition. Data reported by Wu et al.⁶ are also plotted for comparison.

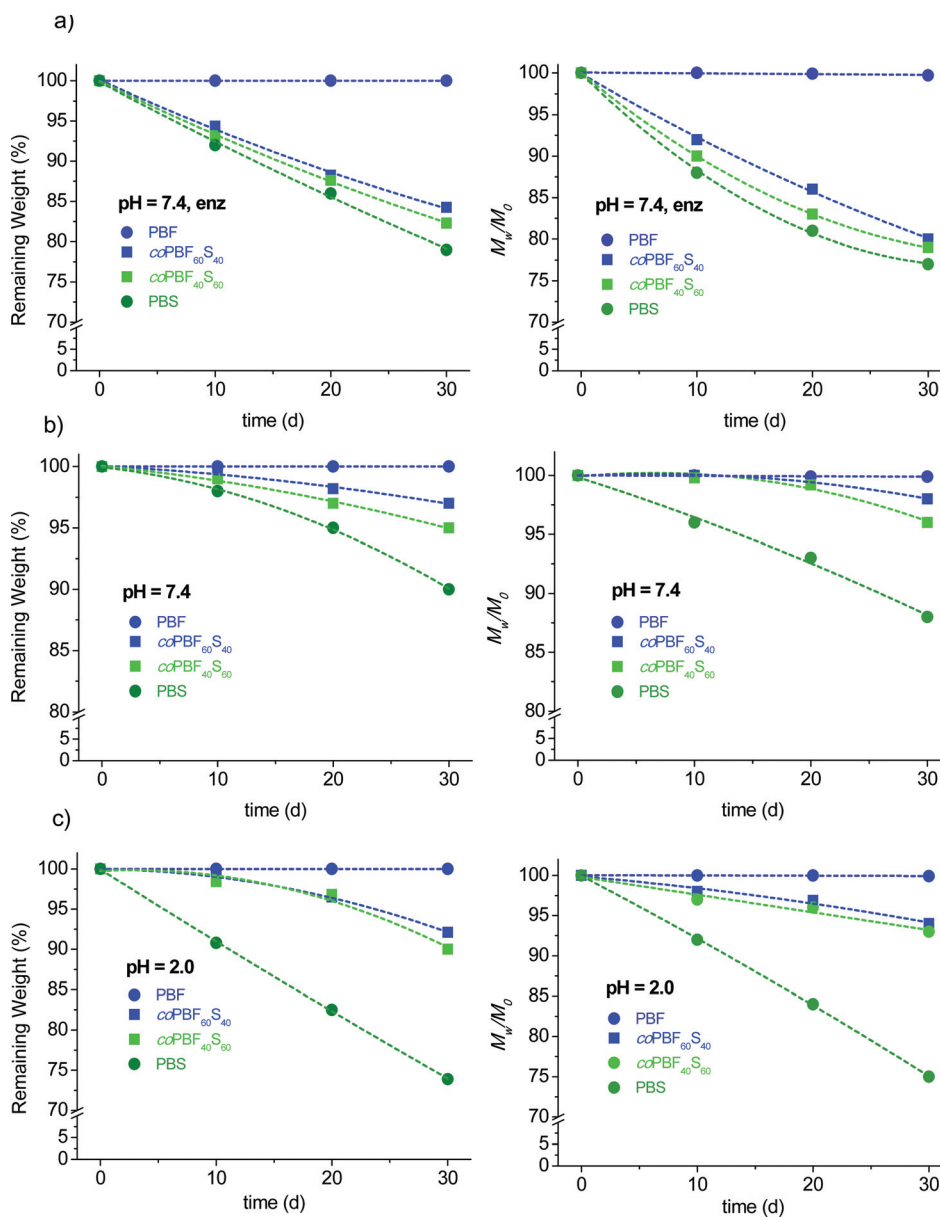


Fig. 10 Enzymatic (a) and hydrolytic degradation (pH 7.4 (b) and pH 2.0 (c)) of coPBF_xS_y . Remaining weight (left) and molecular weight (right) vs. incubation time.



by enzyme catalyzed ROP, which are those presenting higher differences in molecular weight.

3.5. Hydrolytic and enzymatic degradation

It has been reported that PBS is hydrolysable and also biodegradable,^{20,42,43} whereas PBF is known to be highly resistant to hydrolysis and apparently insensitive to biodegradation,⁴⁴ in agreement with the behaviour commonly displayed by aromatic polyesters. It is also known that the degradability of aromatic polyesters may be enhanced by inserting at random aliphatic ester units in the polymer chain. In fact the enzymatic degradation of random poly(butylene furanoate-*co*-adipate) copolyesters has been recently demonstrated, although a minimum content of 50% of adipate units was required to perceive the degradation effects.⁴⁴

In order to evaluate the degradability of PBF as a function of the incorporation of succinate units, the homopolyesters PBF and PBS and the copolyesters *co*PBF₆₀S₄₀ and *co*PBF₄₀S₆₀ were subjected in parallel to degradation assays. Samples were incubated at 37 °C and either pH 2.0 or pH 7.4, in the latter case both in the presence and in the absence of porcine pancreas lipase. The changes taking place in both the sample weight and molecular weight along an incubation period of 30 days were recorded and plotted in Fig. 10. According to what has been reported, PBF remained practically unchanged under all assayed conditions whereas PBS was more or less degraded depending on the pH and whether the enzyme is present or not, showing a maximum of losses at around 25%. As expected, the results obtained for the copolyesters depended on the composition, with degradation being faster as the content of succinate units increased. The degradation rate of the copolyesters was similar at the two assayed pH values, whereas a significant increase was observed in the presence of lipase where both the sample weight and molecular weight losses were around 20%. The conclusion that may be drawn from this preliminary study is that the degradability of PBF can be increased and controlled by the proper adjustment of the amount of succinate units that are incorporated by copolymerization.

4. Conclusions

The results attained in this work demonstrate for the first time that fully bio-based poly(butylene 2,5-furandicarboxylate-*co*-butylene succinate) copolyesters can be successfully synthesized by ROP. Furthermore, the reaction can be either chemically or enzymatically activated by using Sn(Oct)₂ or CALB as the catalyst, respectively. A series of copolyesters covering the whole range of compositions including the two parent homopolyesters could be prepared by chemically catalyzed ROP, whereas only copolyesters containing up to 80 mol% of butylene furanoate units could be prepared by the enzymatic procedure. The ROP method has been proven to be suitable for providing high molecular weight poly(butylene 2,5-furandicarboxylate-*co*-succinate) copolyesters under reaction

conditions milder than those required for melt polycondensation. However, the ROP reactions catalyzed by lipase proceed at significantly higher rates, generating copolyesters of lower molecular weights. The composition of the copolyesters produced by ROP can be readily controlled by adjusting the composition of the feed, and their microstructure is random in all cases. The differences in thermal properties between the two series (obtained by chemically and enzymatically catalyzed ROP) are small and reasonable according to their differences in molecular weights. The preliminary results obtained in this work on biodegradability will contribute to encourage the interest in these copolyesters as fully bio-based materials suitable for short-term applications.

Comparison of our results with those reported by Wu *et al.*⁶ for the same copolyesters produced by melt polycondensation leads us to conclude that the ROP method is a promising option for the synthesis of *co*PBF_{*x*}S_{*y*} copolyesters. However, the true potential of the method in an industrial context will be determined by the accessibility to the cyclic oligomers required for polymerization. The *c*(BS)_{*n*} oligomers are generated as a minor subproduct in the industrial production of PBS by melt polycondensation so that they are readily accessible at large scale. Conversely, the only source today of *c*(BF)_{*n*} is their synthesis in the laboratory, meaning their short-term availability in large amounts is more than unlikely. Nevertheless, the development of PBF chemical recycling processes based on cyclodepolymerization in the not too distant future could be envisaged as a source of these cyclic oligomers. In fact, this method is one of the routes that can be used in the laboratory for their preparation.

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References

- 1 A. Gandini, *Polym. Chem.*, 2010, **1**, 245–251.
- 2 A. Corma, S. Iborra and A. Velty, *Chem. Rev.*, 2006, **107**, 2411–2502.
- 3 C. Vilela, A. F. Sousa, A. Fonseca, A. Serra, J. Cohello, C. Freire and A. Silvestre, *Polym. Chem.*, 2014, **5**, 3119–3141.
- 4 A. F. Sousa, C. Vilela, A. C. Fonseca, M. Matos, C. S. Freire, G. J. M. Gruter, J. F. J. Coelho and A. J. D. Silvestre, *Polym. Chem.*, 2015, **6**, 5961–5983.



- 5 J. Ma, Y. Pang, M. Wang, J. Xu, H. Ma and X. Nie, *J. Mater. Chem.*, 2012, **22**, 3457–3461.
- 6 L. Wu, R. Mincheva, Y. Xu, J. M. Raquez and P. Dubois, *Biomacromolecules*, 2012, **13**, 2973–2981.
- 7 C. H. Wilsens, B. A. Noordover and S. Rastogi, *Polymer*, 2014, **55**, 2432–2439.
- 8 M. Matos, A. F. Sousa, A. Fonseca, C. Freire, J. Coehlo and A. Silvestre, *Macromol. Chem. Phys.*, 2014, **215**, 2175–2184.
- 9 N. Jacquél, R. Saint-Loup, J. P. Pascault, A. Rousseau and F. Fenouillot, *Polymer*, 2015, **59**, 234–242.
- 10 M. Soccio, M. Costa, N. Lotti, M. Gazzano, V. Siracusa, E. Salatelli, P. Manaresi and A. Munari, *Eur. Polym. J.*, 2016, **81**, 397–412.
- 11 G. Z. Papageorgiou, D. G. Papageorgiou, Z. Terzopoulou and D. N. Bikiaris, *Eur. Polym. J.*, 2016, **83**, 202–229.
- 12 J. Xu and G. Bao-Hua, *Biotechnol. J.*, 2010, **5**, 1149–1163.
- 13 I. Bechthold, K. Bretz, S. Kabasci, R. Kopitzky and A. Springer, *Chem. Eng. Technol.*, 2008, **31**, 647–654.
- 14 M. Gigli, M. Fabbri, N. Lotti, R. Gamberini, B. Rimini and A. Munari, *Eur. Polym. J.*, 2016, **75**, 431–460.
- 15 M. Bautista, A. Martínez de Ilarduya, A. Alla and S. Muñoz-Guerra, *Polymers*, 2015, **7**, 1232–1247.
- 16 Q. Liu and X. M. Zhou, *J. Macromol. Sci., Part A: Pure Appl. Chem.*, 2015, **52**, 625–629.
- 17 L. W. Ren, Y. S. Wang, J. Ge, D. N. Lu and Z. Liu, *Macromol. Chem. Phys.*, 2015, **216**, 636–640.
- 18 Y. J. Sun, L. B. Wu, Z. Y. Bu, B. G. Li, N. X. Li and J. M. Dai, *Ind. Eng. Chem. Res.*, 2014, **53**, 10380–10386.
- 19 E. Zakharova, A. Alla, A. Martínez de Ilarduya and S. Muñoz-Guerra, *RSC Adv.*, 2015, **5**, 46395–46404.
- 20 E. Zakharova, C. Lavilla, A. Alla, A. Martínez de Ilarduya and S. Muñoz-Guerra, *Eur. Polym. J.*, 2014, **61**, 263–273.
- 21 R. T. Duan, Q. X. He, X. Dong, D. F. Li, X. L. Wang and Y. Z. Wang, *ACS Sustainable Chem. Eng.*, 2016, **4**, 350–362.
- 22 S. L. Li, F. Wu, Y. Yang, Y. Z. Wang and J. B. Zeng, *Polym. Adv. Technol.*, 2015, **26**, 1003–1013.
- 23 L. C. Tan, Y. W. Chen, W. Zhou, J. C. Wei and S. W. Ye, *J. Appl. Polym. Sci.*, 2011, **121**, 2291–2300.
- 24 M. Lomeli-Rodríguez, M. Martín-Molina, M. Jiménez-Pardo, Z. Nasim-Afzal, S. I. Cauët, T. E. Davies, M. Rivera-Toledo and J. A. Lopez-Sanchez, *J. Polym. Sci., Part A: Polym. Chem.*, 2016, **54**, 2876–2887.
- 25 J. Zhu, J. Cai, W. Xie, P. Chen, M. Gazzano, M. Scandola and R. Gross, *Macromolecules*, 2013, **46**, 796–804.
- 26 A. Gandini and T. M. Lacerda, *Prog. Polym. Sci.*, 2015, **48**, 1–39.
- 27 R. J. I. Knoop, W. Vogelzang, J. Van Haveren and D. S. Van Es, *J. Polym. Sci., Part A: Polym. Chem.*, 2013, **51**, 4191–4199.
- 28 (a) M. Ishii, M. Okazaki, Y. Shibasaki, M. Ueda and T. Teranishi, *Biomacromolecules*, 2001, **2**, 1267–1270; (b) R. Mani, M. Bhattacharya, C. Leriche, L. Nie and S. Bassi, *J. Polym. Sci., Part A: Polym. Chem.*, 2002, **40**, 3232–3239.
- 29 Y. Jiang, A. J. Woortman, G. O. A. van Ekenstein and K. Loos, *Polym. Chem.*, 2015, **6**, 5198–5211.
- 30 (a) S. Sugihara, K. Toshima and S. Matsumura, *Macromol. Rapid Commun.*, 2006, **27**, 203; (b) S. Matsumura, *Macromol. Biosci.*, 2002, **2**, 105–126.
- 31 T. Yagihara and S. Matsumura, *Polymers*, 2012, **4**, 1259–1277.
- 32 H. Ebata, K. Toshima and S. Matsumura, *J. Synth. Org. Chem. Jpn.*, 2008, **66**, 673–382.
- 33 C. W. Lee, K. Masutani and Y. Kimura, *Polymer*, 2014, **55**, 5673–5679.
- 34 D. Pfister, G. Storti, F. Tancini, L. I. Costa and M. Morbidelli, *Macromol. Chem. Phys.*, 2015, **216**, 2141–2146.
- 35 J. C. Morales-Huerta, A. Martínez de Ilarduya and S. Muñoz-Guerra, *Polymer*, 2016, **87**, 148–158.
- 36 J. C. Morales-Huerta, A. Martínez de Ilarduya and S. Muñoz-Guerra, *ACS Sustainable Chem. Eng.*, 2016, **4**, 4965–4973.
- 37 P. Hodge, *Chem. Rev.*, 2014, **114**, 2278–2312.
- 38 D. J. Brunelle, Cyclic Oligomer Chemistry, *J. Polym. Sci., Part A: Polym. Chem.*, 2008, **46**, 1151–1164.
- 39 S. Strandman, J. E. Gautrot and X. X. Zhu, *Polym. Chem.*, 2011, **2**, 791–799.
- 40 A. Kondo, S. Sugihara, M. Kuwahara, K. Toshima and K. Matsumura, *Macromol. Biosci.*, 2008, **8**, 533–539.
- 41 J. C. Randall, in *Polymer Sequence Determination*, New York, 1977, pp. 42–62.
- 42 M. Okada, *Prog. Polym. Sci.*, 2002, **27**, 87–133.
- 43 J. Xu and B.-H. Guo, *Biotechnol. J.*, 2010, **5**, 1149–1163.
- 44 W. Zhou, X. Wang, B. Yang, Y. Xu, W. Zhang, Y. Zhang and J. Ji, *Polym. Degrad. Stab.*, 2013, **98**, 2177–2183.

