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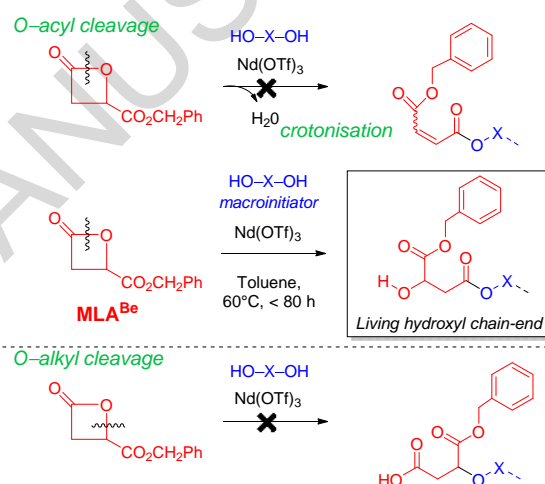
Polyhydroxybutyrate (PHB)-based Triblock Copolymers : Synthesis of Hydrophobic PHB/Poly(benzyl β -malolactonate) and Amphiphilic PHB/Poly(malic acid) Analogues by Ring-Opening Polymerization

Ghislaine Barouti,^a and Sophie M. Guillaume^{*,a}

Polyhydroxyalkanoates (PHAs) are biocompatible and biodegradable polyesters widely used for biomedical applications. Amphiphilic triblock copolymers with PHB hydrophobic segment recently demonstrated attractive advantages such as high colloidal stability and low CMC (Critical Micellar Concentration) values for the elaboration of drug delivery systems. Our approach aims at associating PHB with a fully biodegradable hydrophilic block to access new self-assembled systems with unique potential. Well-defined α,ω -dihydroxy telechelic PHA-based hydrophobic triblock copolymers with tunable segments' length were synthesized by the controlled ring-opening polymerization (ROP) of benzyl β -malolactonate, using polyhydroxybutyrate (PHB) diol/ $\text{Nd}(\text{OTf})_3$ as catalytic system. Remarkably, the reaction proceeds with the selective oxygen-acyl bond cleavage of the β -lactone. The corresponding amphiphilic copolymers were then obtained by hydrogenolysis. These copolymers are the only examples of fully biodegradable PHA-PHB-PHA triblock copolymers. The hydrophilic weight fraction of the copolymers was tuned from 7 to 83% upon modulating the monomer loading thus affording the ability to next access to different self-assembling architectures.

Biodegradable polyesters have been widely studied during the past few decades for various specialty and commodity applications such as in packaging, tissue/bone engineering and drug delivery.¹ Among these, poly(3-hydroxyalkanoate)s (PHAs) such as the ubiquitous poly(3-hydroxybutyrate) (PHB) or the more recently revisited poly(β -malic acid) (PMLA), are extremely attractive due to their additional biocompatibility and biodegradability.²

While natural PHAs can be produced by a number of bacteria, ring-opening polymerization (ROP) of the corresponding cyclic monomers -namely β -lactones such as β -butyrolactone (β -BL) or benzyl β -malolactonate (MLA^{Be}) - is nowadays the most convenient, efficient, versatile and controlled method to synthesize well-defined tailored PHAs with controlled macromolecular characteristics (molar mass predictability, narrow dispersity, chain-end fidelity, microstructure and tacticity).^{3,4} Moreover the high crystallinity of naturally produced isotactic PHB does not allow the high drug loading efficiency (DLE) of hydrophobic drugs, as the result of the low polymer chains mobility which makes the incorporation of the drug more difficult. Typically, ROP of β -lactones may occur through oxygen-alkyl or oxygen-acyl bond cleavage resulting in an inactive carboxylic acid chain-end or in an active hydroxyl terminus, respectively (Scheme 1). Another particularity of this process is the propensity for H_2O abstraction resulting in an inactive crotonate end-capping group.³ Given the challenge in polymerizing these four



Scheme 1. Selectivity issues in the MLA^{Be} cleavage during its ROP initiated by a PHB macrodiol and catalysed by the neodymium triflate salt.

membered ring-strained lactones, the identification of an efficient and selective catalytic system (and to a lesser extent of the operating conditions) is the key point to achieve a good control of the ROP of β -lactones.

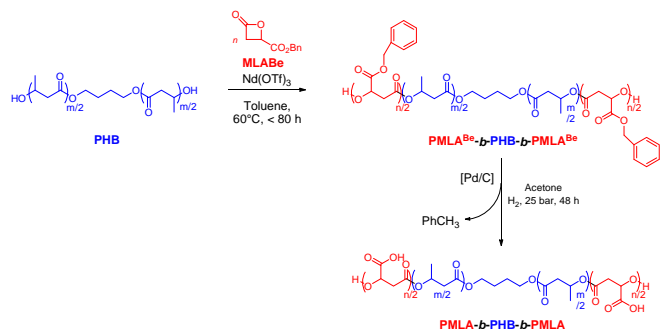
Whereas PHA-based AB diblock copolymers are quite numerous, the analogous ABA triblock copolymers are more limited mostly due to synthetic difficulties. The class of PHB-centered (B block) triblock copolymers includes a few reports with poly(ethylene oxide) A blocks,⁵ with other scarce examples of poly(lactide)^{6a,b} poly(trimethylene carbonate),^{6c} poly(*N*-isopropyl acrylamide),^{6d,e} poly(methacrylate),^{6f} poly(*tert*-butyl acrylate),^{6g} poly(2-(dimethyl-amino)ethyl methacrylate),^{6h} poly(ethyl ethylene phosphate),⁶ⁱ and poly(styrene),^{6f,j} often used within biomaterials.

The association of PHB with $\text{PMLA}^{\text{(Be)}}$ in a triblock copolymer enables some modularity of the properties thanks to the benzyloxycarbonyl (poly(benzyl β -malolactonate), $\text{PMLA}^{\text{(Be)}}$) or carboxylic acid (PMLA) groups resulting in fully hydrophobic PHB/ $\text{PMLA}^{\text{(Be)}}$ or amphiphilic PHB/PMLA copolymers, respectively. In addition, using PHB and PMLA affords fully biodegradable self-assembling systems, whereas in most other PHB-based copolymers,

only a hydrophilic block below a given molar mass is excreted by the renal system. Given that the PHB-*b*-PMLA diblock copolymer afforded promising non-cytotoxic micelles in aqueous solutions,⁷ the parent triblock copolymer PMLA-*b*-PHB-*b*-PMLA thus appears highly attractive due to its potential structuration which may be different as a function of the hydrophilic weight fraction, and which may impart valuable features for drug delivery applications. Indeed, micelles formed from triblock copolymers give more interpenetrated network with favored unimers exchange, thus allowing the possibility to reach a thermodynamic equilibrium.⁸

In this contribution, we report the first synthesis of well-defined hydrophobic PMLA^{Be}-*b*-PHB-*b*-PMLA^{Be} and of their corresponding amphiphilic PMLA-*b*-PHB-*b*-PMLA copolymers, based on the ROP of MLA^{Be} from a PHB-diol macroinitiator and neodymium triflate catalyst (Nd(OTf)₃; OTf = CF₃SO₃), followed by hydrogenolysis of the benzyloxycarbonyl groups under mild conditions (Scheme 2). Along with the previously reported parent diblock copolymers PMLA^(Be)-*b*-PHB,⁷ these copolymers are, to the best of our knowledge, the first such set of analogous diblock and triblock hydrophobic and amphiphilic PHA copolymers.^{2d}

The PMLA^{Be}-*b*-PHB-*b*-PMLA^{Be} triblock copolymers were first synthesized by ROP of MLA^{Be}, using α,ω -dihydroxy telechelic PHB⁹ (Table S1, Fig. S1, ESI[†]) and Nd(OTf)₃ as a new catalytic system (Scheme 2; Table 1), which proceeded through an activated monomer mechanism.^{10,11,12} The monomer conversion determined by ¹H NMR analysis of the crude reaction mixture and the reaction times were in the same range as those reported for the homopolymerization of MLA^{Be} using Nd(OTf)₃/isopropanol or propanediol,^{11,12d} thus highlighting the good efficiency of the dihydroxy functionalized PHB macroinitiator. Varying the MLA^{Be}



cheme 2. Synthesis of PMLA-*b*-PHB-*b*-PMLA triblock copolymer by ROP of MLA^{Be} initiated by Nd(OTf)₃/HO-PHB-OH, followed by hydrogenolysis.

initial loading (5–58 equiv.) and the molar mass of the initial PHB macroinitiator ($M_{n,NMR} = 1400\text{--}6700 \text{ g.mol}^{-1}$, Table S1, ESI[†]), enabled to get a series of PMLA^{Be}-*b*-PHB-*b*-PMLA^{Be} triblock copolymers with various lengths of each segment (PMLA^{Be}/PHB ratios = 13:87 to 90:10).¹³ A set of seven representative hydrophobic triblock copolymers ($M_{n,NMR} = 3300\text{--}11\,500 \text{ g.mol}^{-1}$) was thus obtained in a few grams scale.

The triblock copolymers isolated after a single dialysis were characterized by ¹H NMR spectroscopy in acetone-*d*₆. The samples prepared from the highest MLA^{Be} loading and the longest reaction time revealed that less than 10% of the copolymer chains were end-capped by a carboxylic acid group resulting from the oxygen-alkyl cleavage of the monomer (Table 1, entries 6,7; Fig. S2, Scheme 1, ESI[†]). These -COOH end-functionalized macromolecules were however successfully eliminated by rapid elution of the copolymer sample through a short silica column, as evidenced by the disappearance of the corresponding signal in the ¹H NMR spectra of the recovered samples (*vide infra*, Fig. S5, ESI[†]). No signal characteristic of a crotonate chain-end (C(O)CHCH-C(O)OCH₂Ph, δ 6.85, 5.80 ppm)^{12c-d} was observed in the spectra of the triblock copolymers. Adjusting the operating conditions to a shorter polymerization time thus directly afforded PMLA^{Be}-*b*-PHB-*b*-PMLA^{Be} samples free of macromolecules with carboxylic acid or crotonate termini (Table 1), thereby supporting the selective MLA^{Be} oxygen-acyl bond cleavage.

Table 1. ROP of MLA^{Be} promoted by the Nd(OTf)₃/HO-PHB-OH catalytic system and subsequent hydrogenolysis.

Entry	[MLA ^{Be}] ₀ : [PHB] ₀ : [Nd(OTf) ₃] ₀ ^a	Reaction Time ^b (h)	MLA ^{Be} Conv. ^c (%)	PHB $M_{n,NMR}$ ^d (g.mol ⁻¹)	PMLA ^{Be} - <i>b</i> -PHB - <i>b</i> -PMLA ^{Be} $M_{n,theor}$ (g.mol ⁻¹)	PMLA ^{Be} - <i>b</i> -PHB- <i>b</i> - PMLA ^{Be} $M_{n,NMR}$ ^e (g.mol ⁻¹)	PMLA ^{Be} / PHB NMR molar mass ratio ^g (%)	PMLA ^{Be} - <i>b</i> - PHB - <i>b</i> -PMLA ^{Be} $M_{n,sec}$ ^h (g.mol ⁻¹)	\bar{M}_M ^h	PMLA- <i>b</i> -PHB- <i>b</i> - PMLA M_n ⁱ (g.mol ⁻¹)	PMLA/PHB molar mass ratio ^j (%)
1	5:1:1	7	94	2700	500-2700-500	400-2500-400	24:76	3300	1.29	200-2400-200	14:86
2	6:1:1	10	96	6700	600-6700-600	500-6500-500	13:87	4700	1.33	250-6600-250	7:93
3	8:1:1	12	100	3800	800-3800-800	800-4600-800	26:74	3100	1.26	400-4600-400	15:85
4	12:1:1	40	100	2100	1200-2100-1200	1200-1700-1200	59:41	3200	1.24	600-1700-600	41:59
5	24:1:1	60	100	4900	2500-4900-2500	2200-4900-2200	47:53	5600	1.23	1100-4600-1100	32:68
6	55:1:1	80	100	2100	5700-2100-5700	4100-2500-4100	77:23	3100	1.50	2000-2500-2000	62:38
7	58:1:1	80	98	1400	5900-1400-5900	5200-1100-5200	90:10	3400	1.38	2600-1100-2600	83:17

^a All reactions were run in toluene at 60 °C with an initial concentration of PHB prior to the addition of MLA^{Be} of [PHB]₀ = 0.1 mol.L⁻¹. ^b Reaction times were not necessarily optimized. ^c MLA^{Be} conversion as determined by ¹H NMR spectroscopy of the crude reaction mixture (ESI[†]). ^d Molar mass values of the HO-PHB-OH macroinitiator (not including either the terminal hydrogens or the -O(CH₂)₃O- central moiety) as determined by ¹H NMR spectroscopy of the isolated polymer in acetone-*d*₆ at 25 °C (Table S1, ESI[†]). ^e Theoretical molar mass values of HO-PMLA^{Be}-*b*-PHB-*b*-PMLA^{Be}-OH copolymers (ESI[†]). ^f Molar mass values as determined by ¹H NMR analysis of the isolated block copolymer (ESI[†]). ^g Molar mass ratio of the two blocks of the copolymer as determined by ¹H NMR spectroscopy. ^h Experimental molar mass and dispersity values as determined by SEC analysis in THF at 30 °C vs. polystyrene standards (uncorrected M_n values; ESI[†]). ⁱ Molar mass values of HO-PMLA-*b*-PHB-*b*-PMLA-OH copolymers calculated upon dividing by two the PMLA^{Be} molar mass of the hydrophilic triblock copolymer (ESI[†]), and from the molar mass of PHB as determined by ¹H NMR analysis in acetone-*d*₆ of the isolated amphiphilic copolymer. ^j Molar mass ratio of the PMLA and PHB segments within the copolymer based on the previously determined molar mass value (penultimate column); the first figure refers to the hydrophilic weight fraction.

The purified copolymers were then thoroughly analyzed by NMR spectroscopy as α,ω -dihydroxy telechelic PMLA^{Be}-*b*-PHB-*b*-

PMLA^{Be}. The ¹H NMR spectra of the copolymers showed the distinctive signals corresponding to both BL and MLA^{Be} repeating

units and the central tetramethylene moiety, as illustrated Fig. 1 (Fig. S3-S5, ESI⁺). The good resolution of these latter methylene resonances enabled their fairly reliable integration from which both the PHB and PMLA^{Be} segments molar mass values ($M_{n,NMR}$) could be determined. These data were in good agreement with the ones calculated from the monomer conversion and from the initial feed of MLA^{Be} and of the initiator ($M_{n,theo}$), as reported in Table 1. The corresponding $^{13}\text{C}\{^1\text{H}\}$ J-MOD, $^1\text{H}-^{13}\text{C}$ HMBC and $^1\text{H}-^{13}\text{C}$ HSQC NMR spectra similarly evidenced and confirmed the presence of PHB and PMLA^{Be} units, and of the tetramethylene central moiety (Fig. S6-S8, ESI⁺). Furthermore, DOSY NMR experiments are being more and more implemented to evidence the formation of block copolymers and to assess their purity.^{7,14} DOSY NMR analyses of the PMLA^{Be}₂₂₀₀-*b*-PHB₄₉₀₀-*b*-PMLA^{Be}₂₂₀₀ triblock copolymer showed a single diffusion coefficient for all the signals, distinct from the two coefficients observed for a mixture of the PHB-diol macroinitiator and of a PMLA^{Be} homopolymer with comparable molar mass to the triblock composition (Fig. 2). These results suggested the absence of contamination of the triblock copolymer by any homopolymer, and the presence of a single macromolecular species in the sample.

SEC analysis in THF of the purified triblock copolymer samples typically showed a trace shifted to a higher molar mass value as compared to that of the corresponding PHB macroinitiator (Fig. S9, ESI⁺). No residual PHB macroinitiator was therein observed thus supporting the high efficiency of the macro-diol to initiate the ROP of MLA^{Be}, and that all polymer chains were initiated by HO-PHB-OH. Monomodal chromatograms with narrow dispersity values ($D_M = 1.23-1.50$, ESI⁺), along with the good agreement of $M_{n,theo}$ and $M_{n,NMR}$ values (*vide supra*, Table 1), suggested the good control of the polymerization.¹⁶

The amphiphilic α,ω -dihydroxy telechelic triblock copolymers HO-PMLA-*b*-PHB-*b*-PMLA-OH were next obtained upon hydrogenolysis of the parent hydrophobic copolymers HO-PMLA^{Be}-*b*-PHB-*b*-PMLA^{Be}-OH by using a heterogeneous Pd/C catalyst under

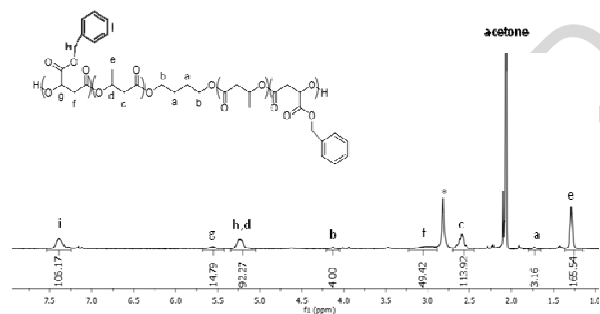


Fig. 1. ^1H NMR (400 MHz, acetone- d_6 , 23 °C) spectrum of HO-PMLA^{Be}₂₂₀₀-*b*-PHB₄₉₀₀-*b*-PMLA^{Be}₂₂₀₀-OH synthesized by ROP of MLA^{Be} from Nd(OTf)₃/HO-PHB₄₉₀₀-OH (Table 1, entry 5) (* marker stands for residual water).¹⁵

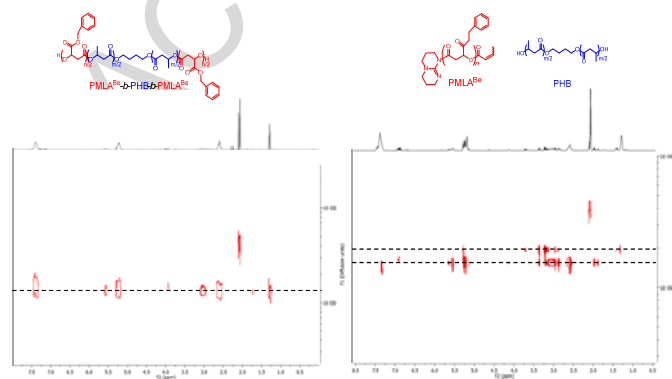


Fig. 2. DOSY NMR (400 MHz, acetone- d_6 , 23 °C) spectrum of HO-PMLA^{Be}₂₂₀₀-*b*-PHB₄₉₀₀-*b*-PMLA^{Be}₂₂₀₀-OH ($D = 127.10 \cdot 10^{-11} \text{ m}^2 \cdot \text{s}^{-1}$) synthesised by ROP of MLA^{Be} from Nd(OTf)₃/HO-PHB₄₉₀₀-OH (Table 1, entry 5) and of a mixture of HO-PMLA^{Be}₄₀₀₀-crotonate ($D = 154.10 \cdot 10^{-11} \text{ m}^2 \cdot \text{s}^{-1}$) and HO-PHB₅₀₀₀-OH ($D = 191.10 \cdot 10^{-11} \text{ m}^2 \cdot \text{s}^{-1}$).

mild conditions (Scheme 2). This enabled the cleavage of the pendant benzyloxycarbonyl moieties of the MLA^{Be} repeating units without alteration of the polymer backbone, a well-known abstraction method suitable for polyesters (Scheme 2).^{2c,7,14a,18} The amphiphilic PMLA-*b*-PHB-*b*-PMLA copolymers were then recovered as a viscous colorless oil after dialysis (Table 1).

In order to characterize these amphiphilic triblock copolymers by NMR spectroscopy, a good solvent of both segments needed to be identified. However, no common organic solvent (acetone, acetone/ trifluoroacetic acid (TFA), acetone/water, THF, DMSO, DMF, CHCl₃, CH₂Cl₂) was found suitable to simultaneously solubilize both blocks. The amphiphilic character of the copolymer and the triblock architecture most likely enhanced the ability of the copolymer to somehow self-organize, thus compromising its solubility and characterization. Nevertheless, analysis in acetone- d_6 first evidenced the total disappearance of the benzylic signals of the pending benzyl protecting groups, thus supporting the complete deprotection of the PMLA^{Be} segment into the PMLA block, and further showed a basically unchanged molar mass value of the PHB block after hydrogenolysis, as expected (Fig. S10, ESI⁺). Addition of some TFA to the above mentioned solvents (2–50%, v/v), as successfully enabling the complete dissolution of the related PMLA-*b*-PHB diblock copolymers,⁷ also failed to simultaneously solubilize both PHA segments of the triblock copolymers. The addition of 2% of TFA (v/v) to acetone- d_6 only resulted in the cleavage of the hydrogen bonds thus allowing, in particular, to clearly observe the resonance of the PMLA pending -COOH groups (Fig. 3, S11-S13, ESI⁺). Yet, this did not enable to fully dissolve the PMLA segment thus precluding the evaluation of its molar mass by ^1H NMR spectroscopy. Nevertheless, the molar mass was estimated to theoretically amount to half the molar mass of PMLA^{Be}, as the result of the depletion of the benzyloxy groups (Be corresponds to half the molecular weight of MLA^{Be}) (Table 1). A series of amphiphilic triblock copolymers featuring various hydrophilic PMLA fractions (*f ca.* 7–83%) was thus successfully isolated.

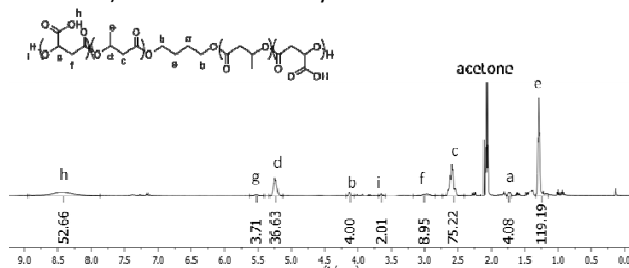


Fig. 3. ^1H NMR (400 MHz, acetone- d_6 /TFA (98:02, v/v), 23 °C) spectrum of HO-PMLA-*b*-PHB-*b*-PMLA-OH obtained upon hydrogenolysis of HO-PMLA^{Be}₂₂₀₀-*b*-PHB₄₉₀₀-*b*-PMLA^{Be}₂₂₀₀-OH (Table 1, entry 5). The signals of the PHB and PMLA block do not suitably integrate as the result of the partial solubility of the triblock copolymer in this solvent.

The corresponding $^{13}\text{C}\{^1\text{H}\}$ J-MOD NMR spectra of the PMLA-*b*-PHB-*b*-PMLA copolymers in acetone- d_6 /TFA similarly evidenced the characteristic signals of PMLA and PHB segments and of the tetramethylene central moiety (Fig. S14, ESI⁺). Moreover, DOSY NMR analyses of the amphiphilic copolymers revealed a single diffusion coefficient ($D = 102.10 \cdot 10^{-11} \text{ m}^2 \cdot \text{s}^{-1}$), different from that of the protected copolymer (Fig. 2 vs S15, ESI⁺), thus supporting that the sample fits well with a single-population model.¹⁹ All these NMR analyses showed that the hydrophobic triblock copolymers were smoothly chemically modified into their parent amphiphilic copolymers without fragmentation of the backbone and blocks.

Furthermore, the surprisingly lower D value of the lower molar mass deprotected triblock copolymer as compared to the protected precursor, hinted that these triblock copolymers behaved distinctively than the analogous hydrophobic/amphiphilic diblock copolymers.⁷ This also suggested that not only the hydrophilic weight fraction f of an amphiphilic copolymer is important to understand its physico-chemical behavior, but also that its architecture as a diblock or triblock copolymer also dictates to some extent its behavior in solution. The copolymer chemical composition-architecture-solution behavior relationship is thus a significant notion to investigate to better understand these PHAs triblock copolymers.

The thermal properties of PMLA^{Be}-*b*-PHB-*b*-PMLA^{Be} and PMLA-*b*-PHB-*b*-PMLA were investigated by differential scanning calorimetry (DSC) and thermogravimetric analysis (TGA) (Fig. S17, S18, ESI[†]). The hydrophobic triblock copolymer (Table 1, entry 7) showed two glass transitions temperatures ($T_g = +1, +26$ °C, corresponding to the PHB and PMLA^{Be} blocks, respectively (ESI[†]), a behavior similarly observed for PHB-*b*-PMLA^{Be} copolymers.⁷ The thermal degradation of the hydrophobic copolymers was found to be much slower and to occur at a higher temperature than that of the corresponding amphiphilic copolymers (f 41, 83; Table 1, entries 4, 7), respectively. Both sets of triblock copolymers exhibited a stepwise degradation profile corresponding to the degradation of first the PMLA^{Be} segment and then the PHB segment, as similarly observed for the PHB-*b*-PMLA^{Be} copolymers.⁷ (ESI[†]). These thermal behaviors again highlighted the significant effect of the chemical composition/modification of the segment(s) of the block copolymers.

In summary, well-defined analogous hydrophobic and amphiphilic PHA-based triblock copolymers were successfully synthesized from commercially available reagents (except for MLA^{Be}), by the efficient and controlled ROP of MLA^{Be} using a PHB diol/Nd(OTf)₃ catalytic system nicely mediating the selective oxygen-acyl bond cleavage of the β -lactone, followed by hydrogenolysis. These copolymers represent, to our knowledge, the only examples of PHA-PHB-PHA triblock copolymers. This strategy could be extended to the design of other PHA-based ABA and BAB triblock copolymers. These novel α,ω -dihydroxy telechelic triblock PHA copolymers are highly valuable. Indeed, they may also serve as macroinitiators towards the synthesis of other types of block copolymers. Also, thanks to the pending -COOH moieties, they provide anchoring sites for biological molecules of interest towards conjugated polymers. Finally, through their amphiphilicity, they may be valorized as nanostructured entities for drug delivery systems. Investigations of the self-assembling behavior of the PMLA-*b*-PHB-*b*-PMLA copolymers revealed unique properties in relation to the hydrophilic weight fraction. Detailed results and outcomes along these lines will be reported in due course.

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- intermolecular (reshuffling) and intramolecular (back-biting) transesterification reactions and/or other chain transfer or termination reactions), and/or an initiation somewhat slower than the propagation. Furthermore, these dispersity values are in the range of those typically obtained for PMLA^{Be} homopolymers and PMLA^{Be}/PHB copolymers.^{11,12,18a,c}
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 16. Note that, the experimental molar mass values assessed by SEC analyses in THF ($M_{n,SEC}$, Table 1) also fairly matched the PMLA^{Be}-*b*-PHB-*b*-PMLA^{Be} $M_{n,NMR}$ and $M_{n,theo}$ data for the copolymers featuring short PMLA^{Be} blocks. Indeed, $M_{n,SEC}$ of PMLA^{Be}-enriched copolymers (*ca.* > 15 molar mass%) remained lower than the expected values, possibly reflecting the adsorption of these copolymers onto the columns. Such a behaviour was previously observed for related PMLA^{Be} and PMLA^{Be}/PHB (co)polymers.^{7,11,12c-d,14a,17}
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