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Supporting for

Synthesis and properties of ABA-triblock copolymers from polyesters A-blocks and polyacetals B-blocks with enhanced ability to degradation

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Assay of Sulphur content

0.2 g of copolymers were placed into glass vessels, then 65% nitric acid (3 mL) and 36% hydrochloric acid (1 mL) spectrally pure were added. Samples were pre-mineralized at 25 °C within 25 min. by using an ultrasonic bath. Then, the samples were mineralized in the UltraWave closed microwave system in two steps, as presented in the Table 1.

Table S1. Conditions of mineralization

Step	Time [min.]	T1 [⁰ C]	T2 [⁰ C]	Pressure [Ba]	Power [W]
1	25	240	70	110	1500
2	10	240	70	110	1500
2	10	240	/0	110	1300

After mineralization, the samples were transferred to 25 mL volumetric flasks and filled with deionized water and characterized. The Sulphur content was determined to be 932 ppm.

Table S2. The diffusion coefficient for signals belonging to the PDXL $_{7000}$ and PLA $_{7500}$ -PDXL $_{7000}$ -PLA $_{7500}$ block copolymer

PDXL_{7000}					
Signal [ppm]	Difusion coefficiency				
4.75	1.26·10-6				
3.70	1.26·10-6				
PLA ₁₀₀₀₀					
5.17	1.32·10-6				
1.60	1.34·10 ⁻⁶				
PLA ₇₅₀₀ -PDXL ₇₀₀₀ -PLA ₇₅₀₀					
5.12	9.95·10 ⁻⁷				
4.75	1.02·10-6				
4.40-4.28	1.02·10 ⁻⁶				
3.70	$9.99 \cdot 10^{-7}$				
1.60	1.01 · 10-6				

Table S3. Thermal ABA copolymers (the

properties of selected

second heating cycle at a rate of 10 °C/min in a DSC analysis)

Copolymer	Ester content	DSC				TGA	
	mol %	T _g [°C]	T _m PDXL [°C]	T _m polyester [°C]	ΔH_{m} [J/g]	T ₁ [°C]	T ₂ [°C]
PLA-PDXL ₇₀₀₀ -PLA 1	26	-41				227	308
PLA-PDXL ₇₀₀₀ -PLA 2	35	-26				229	299
PLA-PDXL ₇₀₀₀ -PLA 3	44	-13		123	12.78	239	298
PLA-PDXL ₇₀₀₀ -PLA 4	53	3		145	23.51	239	308
PLA-PDXL ₇₀₀₀ -PLA 5	60	3		150	27.24	243	324
PCL-PDXL ₇₀₀₀ -PCL 1	42	-63	31	53	68.34	305	355
PCL-PDXL ₇₀₀₀ -PCL 2	65	-65	24	51	62.74	302	371

 $T_{\rm g}$, $T_{\rm m}$ - temperature of glass transition and melting

 $\Delta H_{\rm m}$ - polyester enthalpy of melting

 T_1 , T_2 - temperature of the maximum decomposition

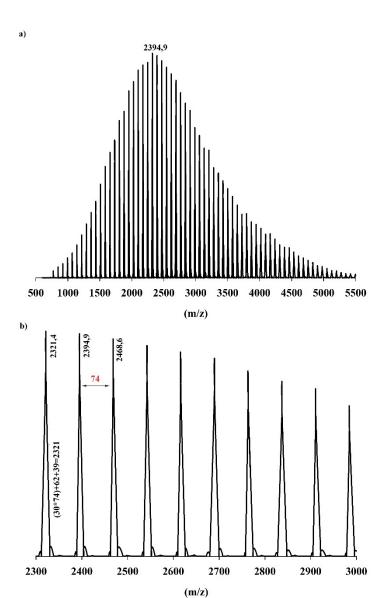


Figure S1. MALDI TOF of purified PDXL (M_{nNMR} = 2400 g/mol)

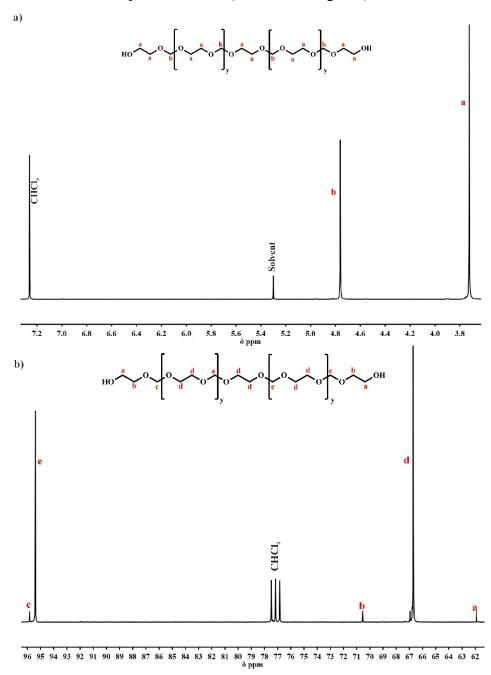


Figure S2. The ¹H and ¹³C NMR spectra (400 MHz, CDCl₃) of purified PDXL₇₀₀₀

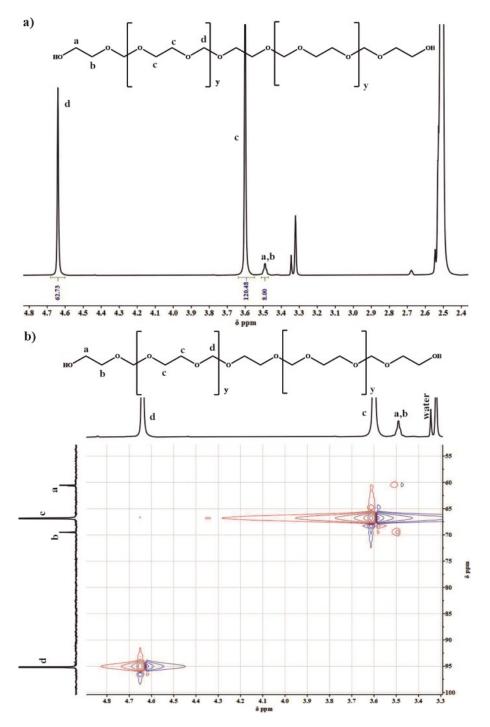


Figure S3. ^1H NMR spectrum, and 2D HSQC NMR (400 MHz, DMSO-d₆) of PDXL₂₄₀₀ diol

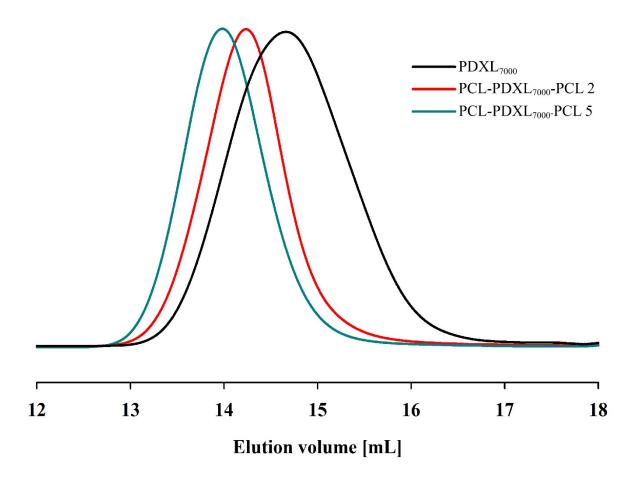
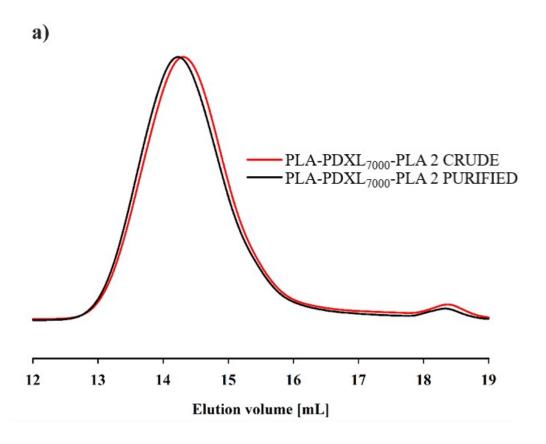


Figure S4. SEC trace of polyacetal macroinitiator HO-PDXL $_{7000}$ -OH and prepared purified PCL-PDXL $_{7000}$ -PCL copolymers



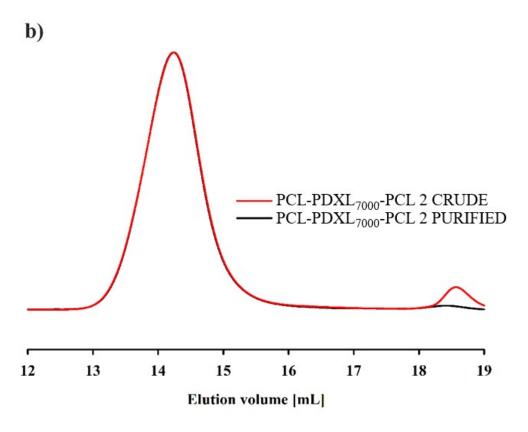


Figure S5. The SEC traces of crude and purified a) PLA-PDXL₇₀₀₀-PLA copolymer and b) PCL-PDXL₇₀₀₀-PCL copolymer

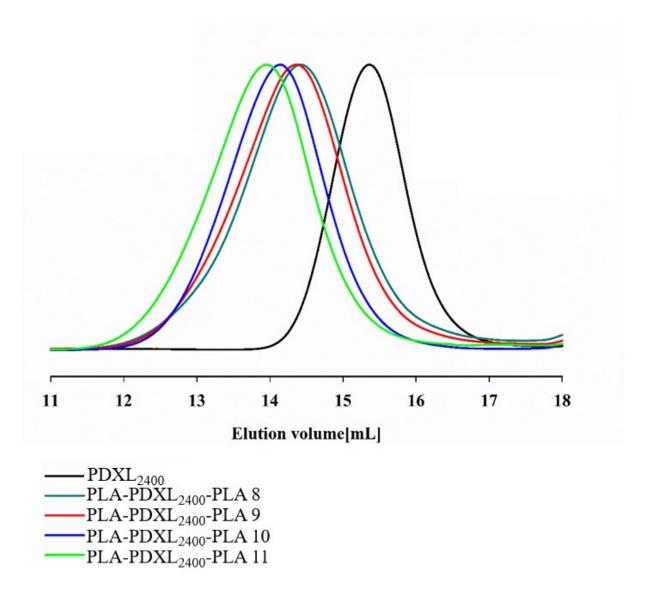


Figure S6. SEC curves of polyacetal macroinitiator HO-PDXL $_{2400}$ -OH and resulting purified PLA-PDXL $_{2400}$ -PLA copolymers.

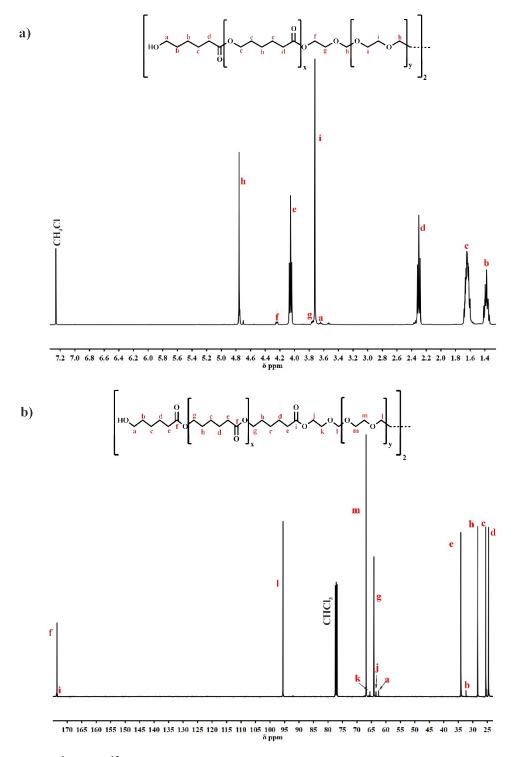


Figure S7. The 1 H and 13 C NMR spectra (400 MHz, CDCl₃) of purified PCL-PDXL₇₀₀₀-PCL copolymer.

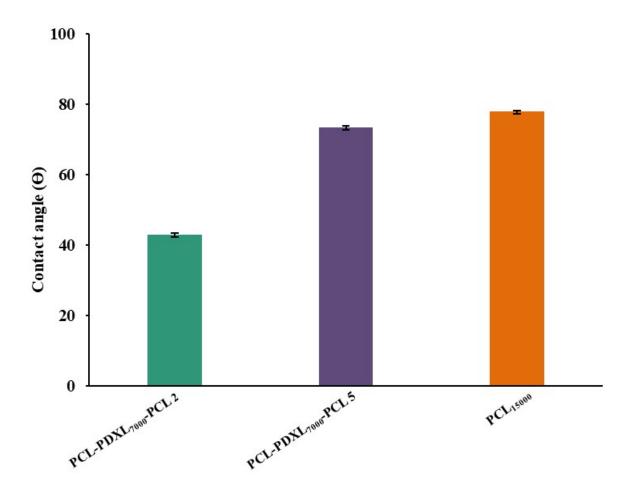


Figure S8. Assessment of mean static contact angles of PCL-PDXL $_{7000}$ -PCL block copolymers in comparison to pure PDXL $_{7000}$ and PCL $_{14000}$.

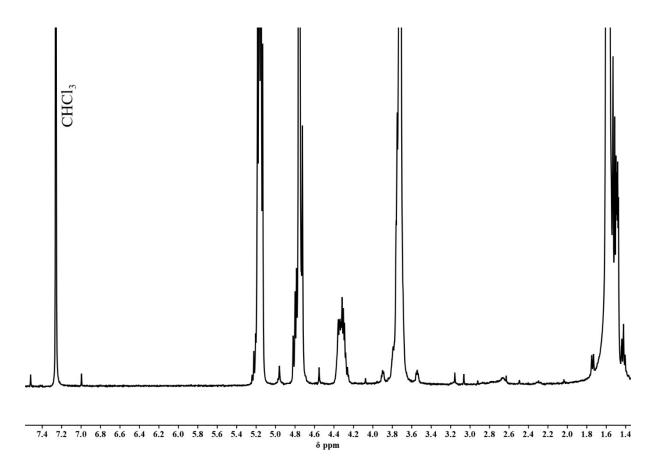


Figure S9. The ¹H NMR of the films for the WCA measurements (CDCl₃, 400 MHz)

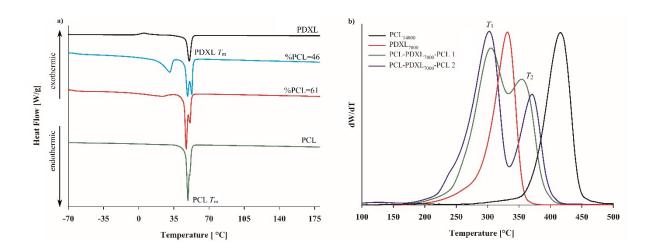


Figure S10. a)The DSC thermograms registered during the second scan and b) TGA curves for starting HO-PDXL₇₀₀₀-OH diol, PCL₁₄₀₀₀, and PCL-PDXL₇₀₀₀-PCL. For DSC analysis samples were heated at 10 °C/min and cooled at 10 °C/min, whereas for TGA measurement samples were heated at 20 °C/min.

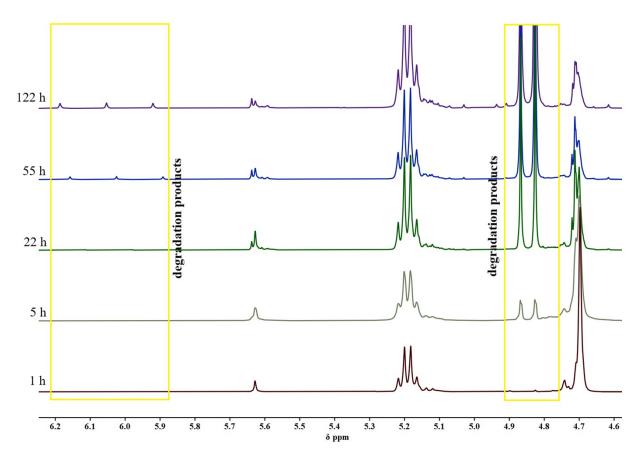


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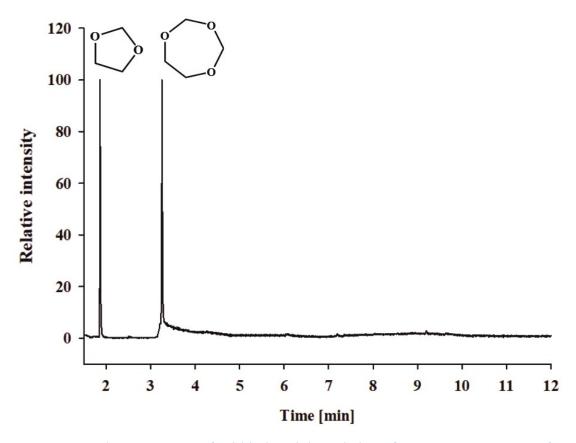


Figure S12. The GC spectra of acid induced degradation of PLA-PDXL₇₀₀₀-PLA after 122h