Supporting Information for

Synthesis and characterization of poly(ethylene terephthalate-

co-1,4-cyclohexanedimethylene terephtlatate)- block-

poly(tetramethylene oxide) copolymers

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Characterization method.

The ¹H-¹³C gHSQC (gradient enhanced heteronuclear single quantum coherence) 2D-NMR experiment was performed with a standard Varian pulse sequence with 90° pulse widths of 9.1 and 8.6 μ s for ¹H and ¹³C, respectively. The ¹H dimension had a spectral width of 9.6 kHz, and the ¹³C dimension had a spectral width of 25.6 kHz, acquisition time of 0.15 s, 1.0 s relaxation delay; 16 transients were averaged for each of 256 increments using the States method of phase sensitive detection in *f*₁. Delays were optimized depending on the coupling interests. The data were zero-filled to 4096 × 4096 and weighted with sine bell and shifted sine bell functions prior to Fourier transformation.

The ¹H-¹H gCOSY (gradient correlation spectroscopy) 2D NMR experiments were performed with the standard Agilent sequence. This was done with a 9.6 kHz spectral width, 0.15 s acquisition time, 1 s relaxation delay, and 9.1 μ s 90° pulse width; a total of 8 transients were averaged for each 256 increments using the States method of phase sensitive detection in f_1 . Processing was done with shifted sine bell weighting functions in both dimensions and zero-filling to a 1024 × 1024 data matrix prior to Fourier transformation.

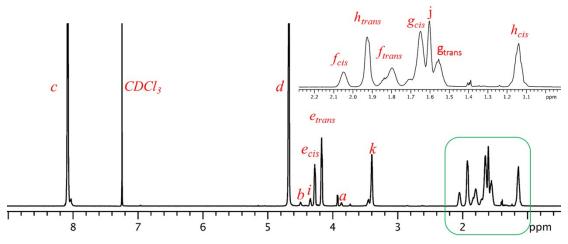


Figure S1. ¹H quantitative NMR spectra of PEE 90/10 sample. Solvent: CDCl₃.

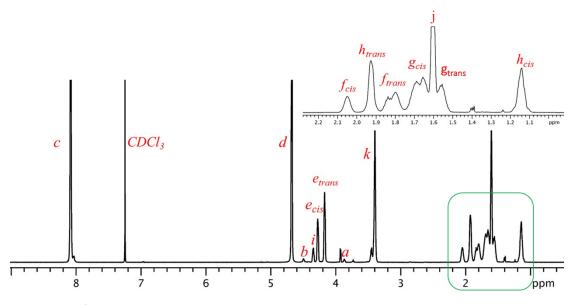


Figure S2. ¹H quantitative NMR spectra of PEE 80/20 sample. Solvent: CDCl₃.

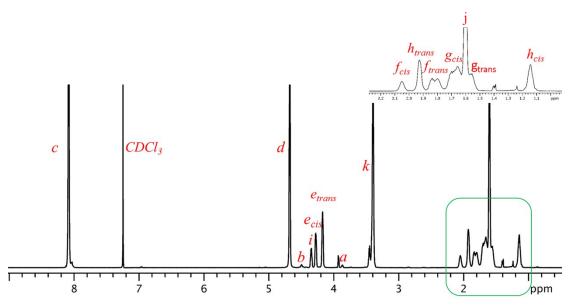


Figure S3. ¹H quantitative NMR spectra of PEE 70/30 sample. Solvent: CDCl₃.

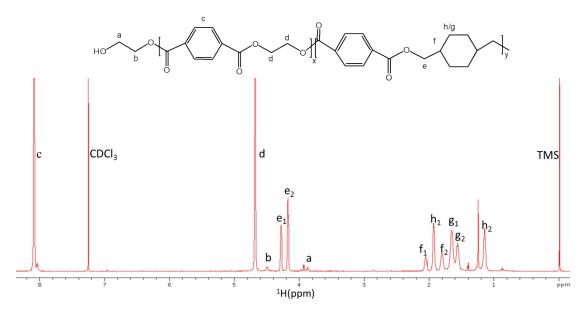


Figure S4. ¹H quantitative NMR spectra of PETG sample. Solvent: CDCl₃.

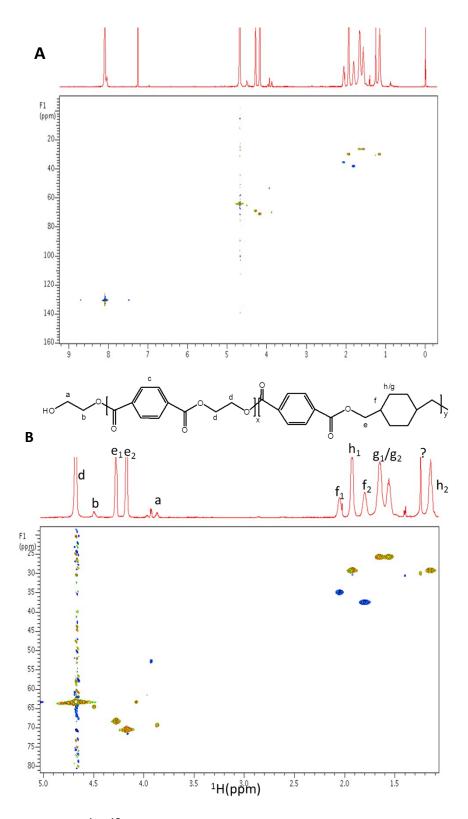


Figure S5. ${}^{1}H{}^{-13}C$ gHSQC NMR full spectrum (A) and enlarged selected region (B) of PETG sample. Solvent: CDCl₃.