

Preparation and property investigation of chain end functionalized *cis*-1,4 polybutadienes via de-polymerization and cross metathesis of *cis*-1,4 polybutadienes

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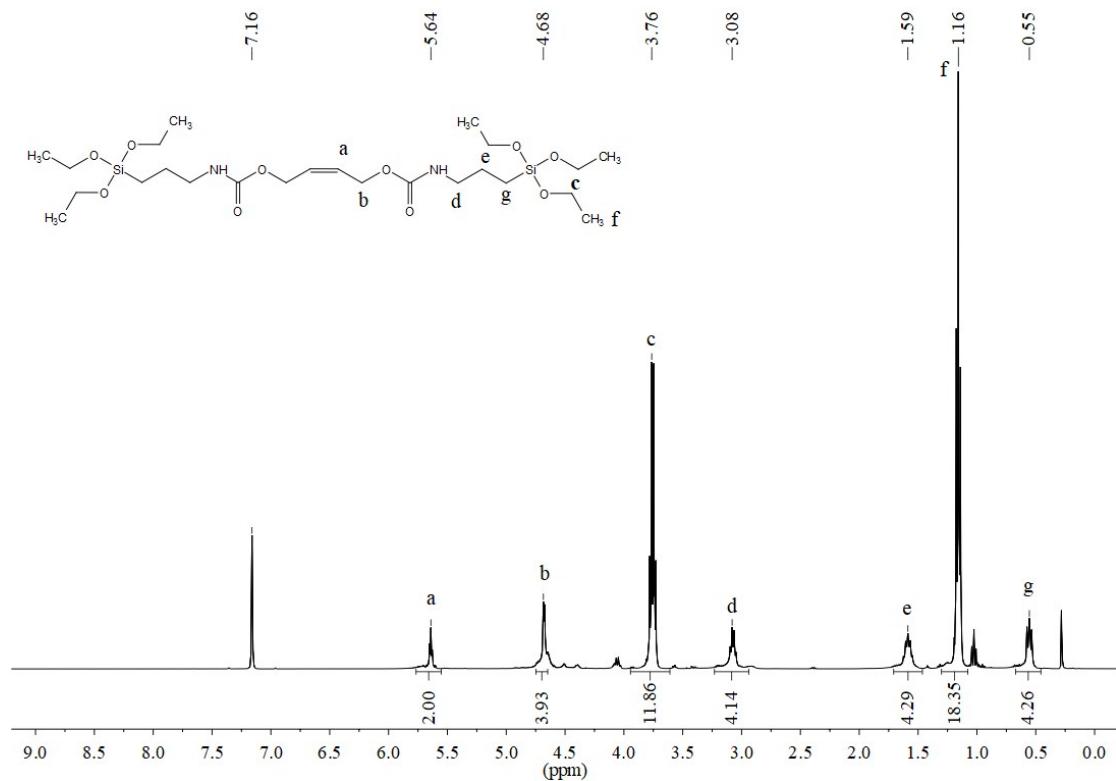


Figure 1S ^1H NMR spectra of CTA-Si(OEt)₃

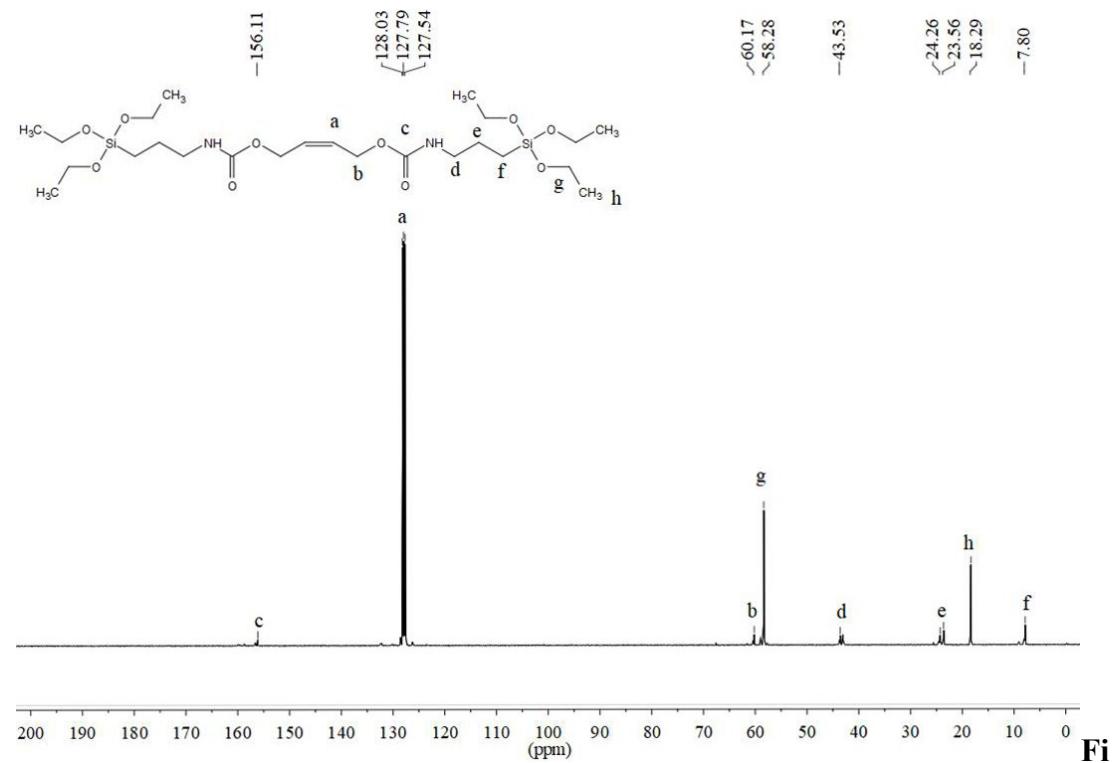


Figure 2S ^{13}C NMR spectra of CTA-Si(OEt)₃

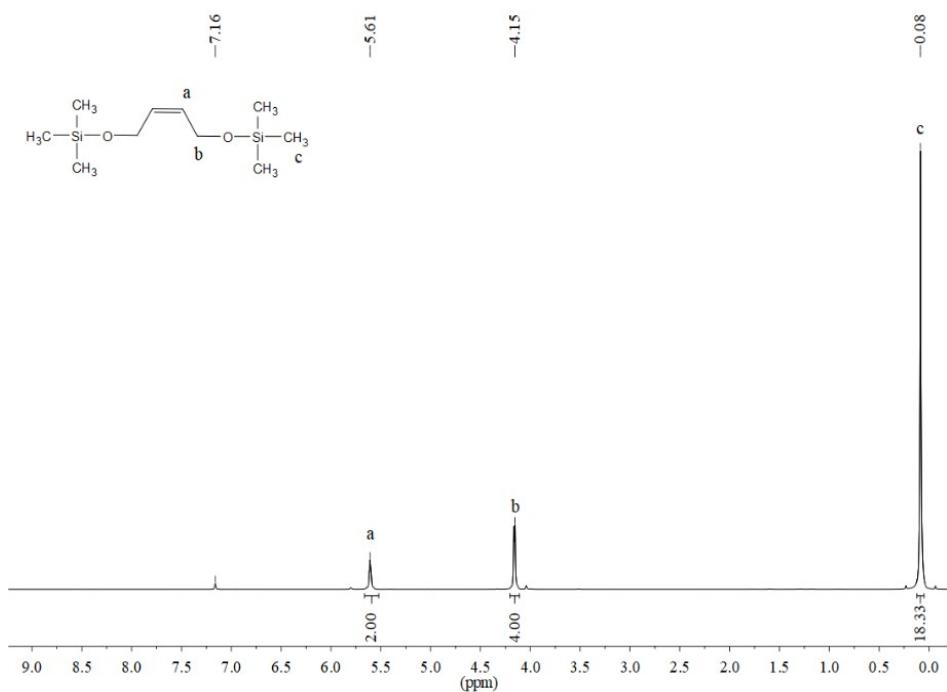


Figure 3S ¹H NMR spectra of CTA-SiMe₃

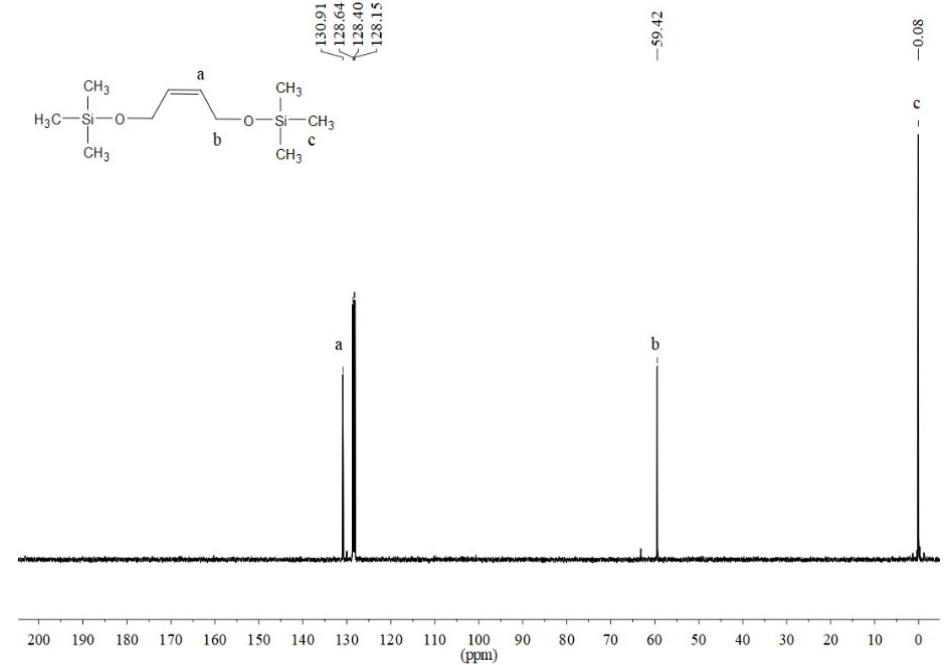


Figure 4S ¹³C NMR spectra of CTA-SiMe₃

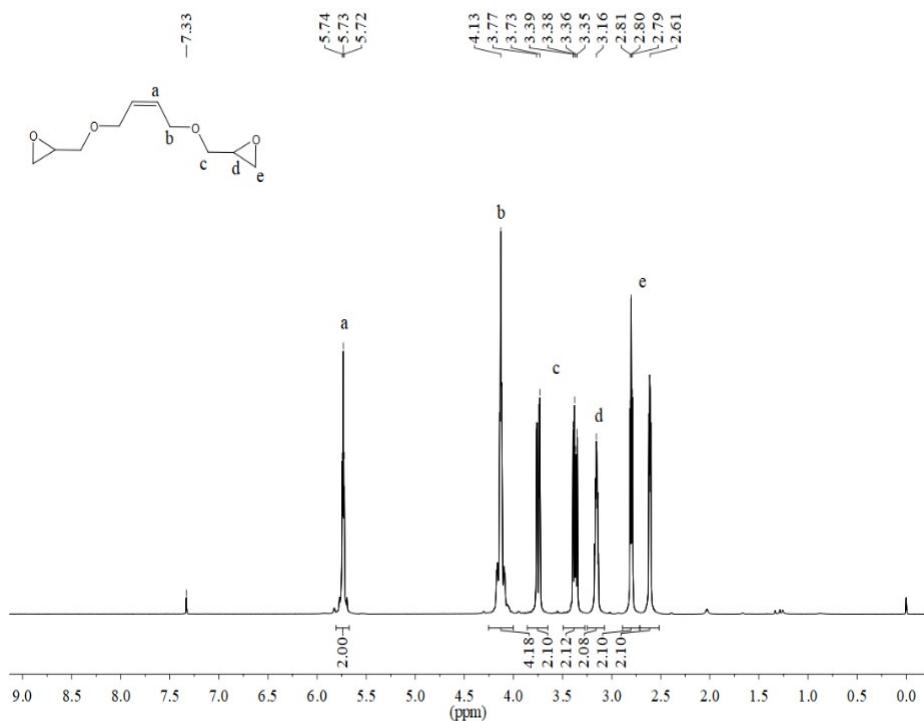


Figure 5S ^1H NMR spectra of CTA-PO

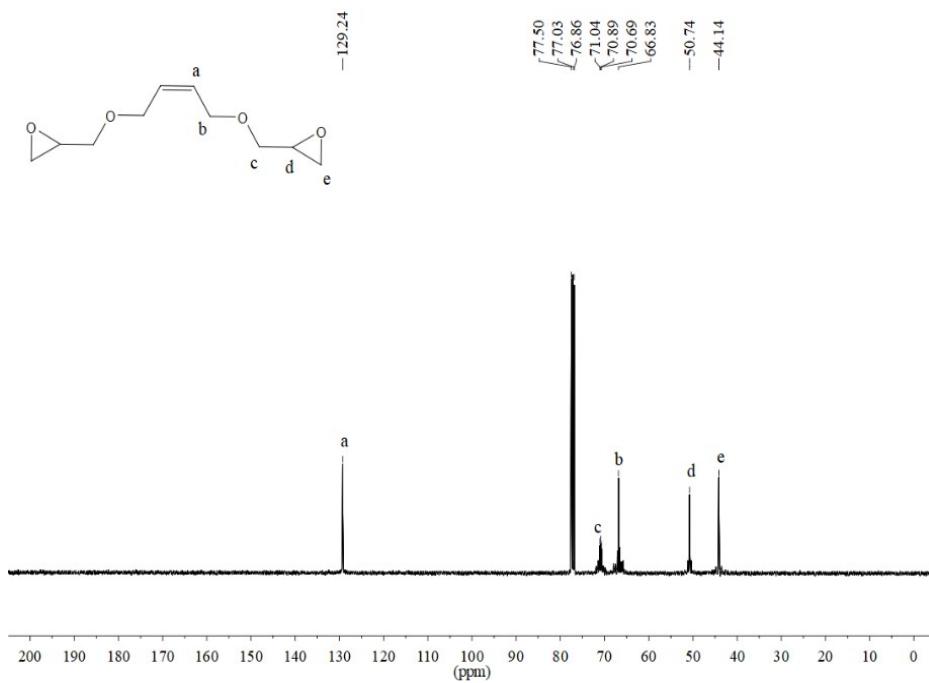


Figure 6S ^{13}C NMR spectra of CTA-PO

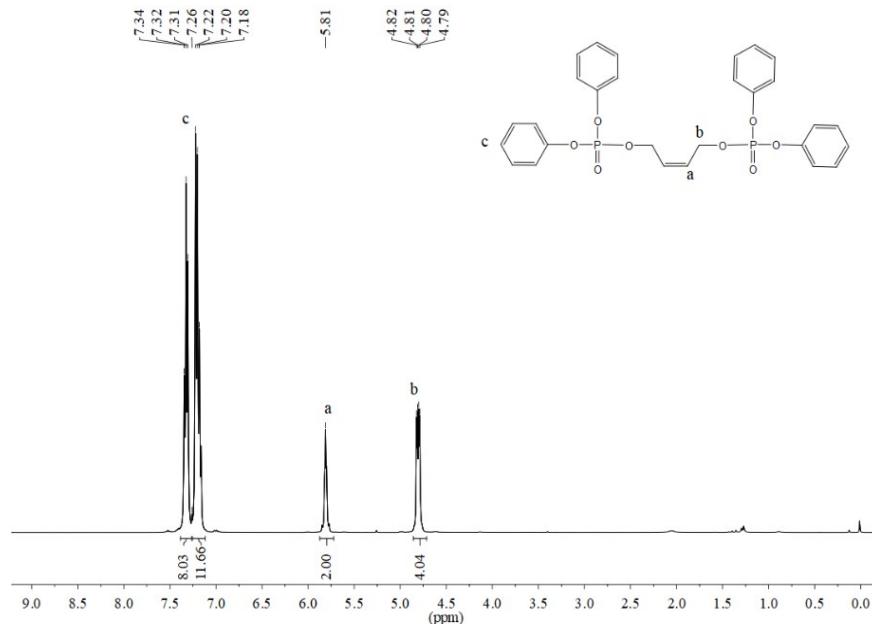


Figure 7S ^1H NMR spectra of CTA-Phosphate

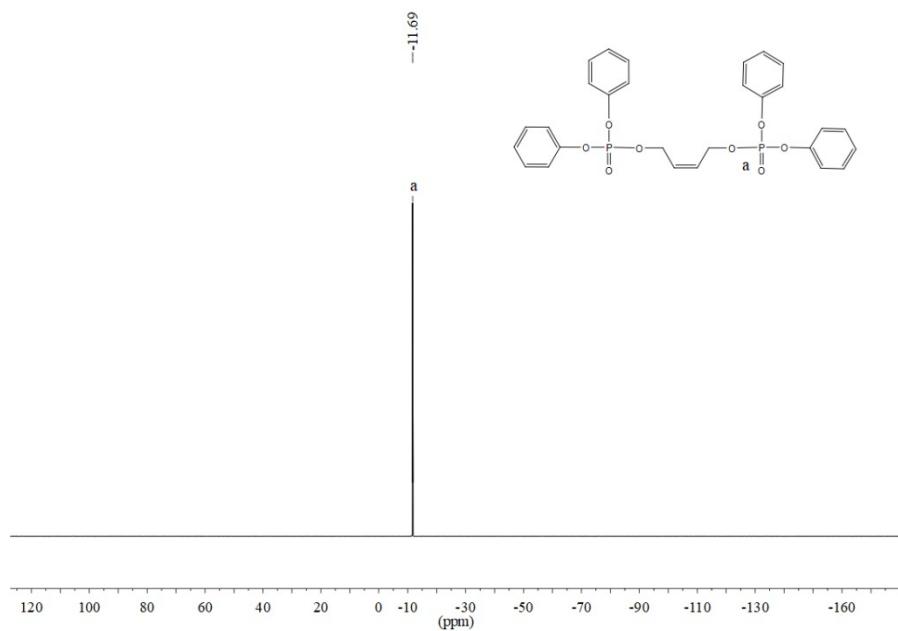


Figure 8S ^{31}P NMR spectra of CTA-Phosphate

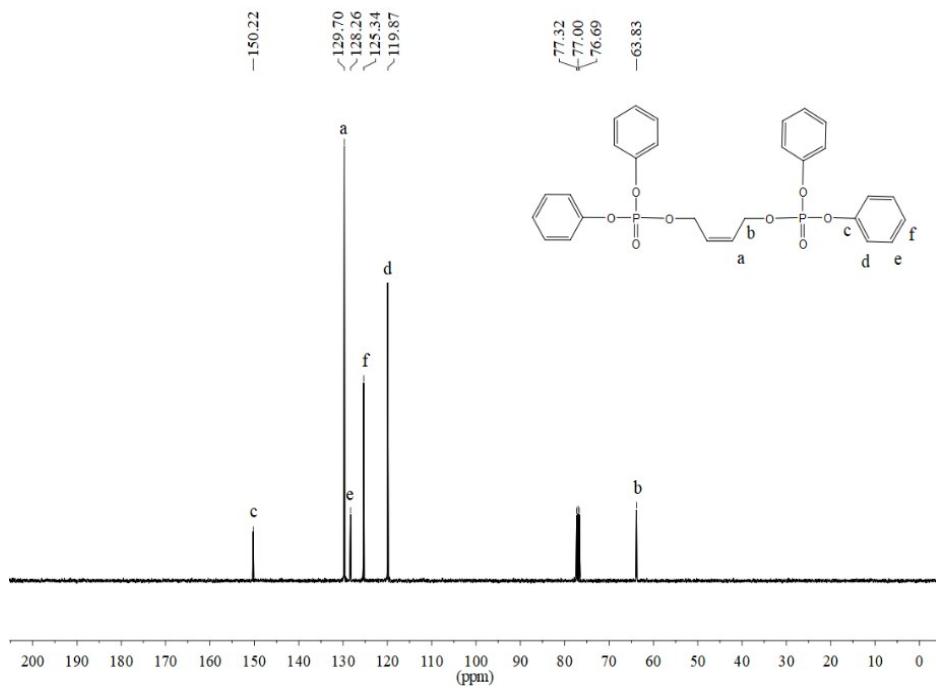


Figure 9S ^{13}C NMR spectra of CTA-Phosphate

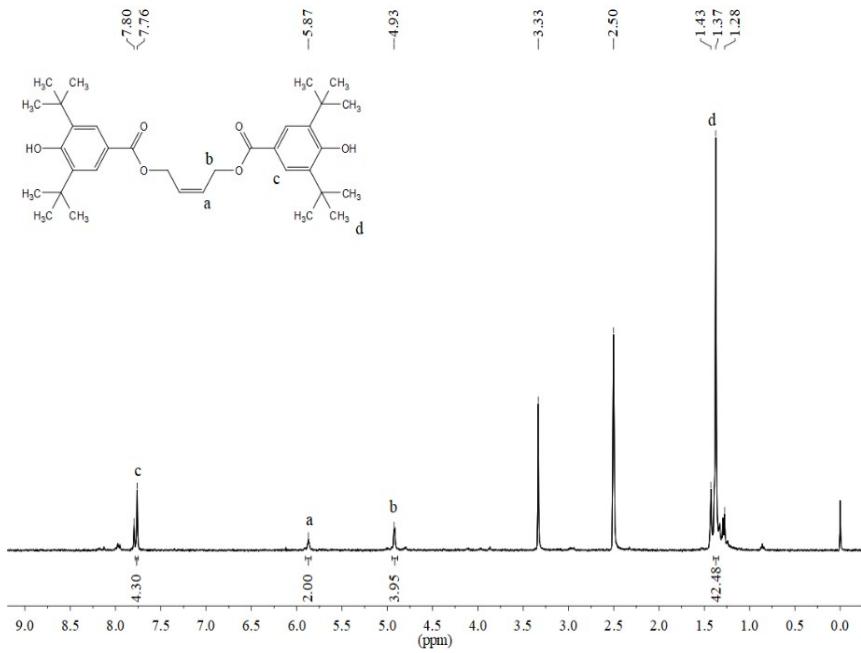


Figure 10S ^1H NMR spectra of CTA-Phenol

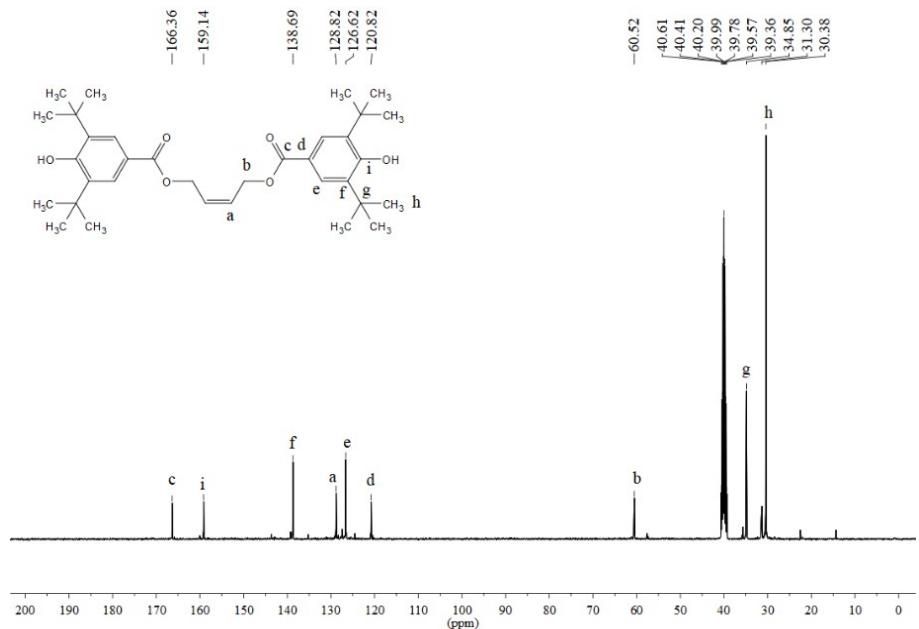


Figure 11S ^{13}C NMR spectra of CTA-Phenol

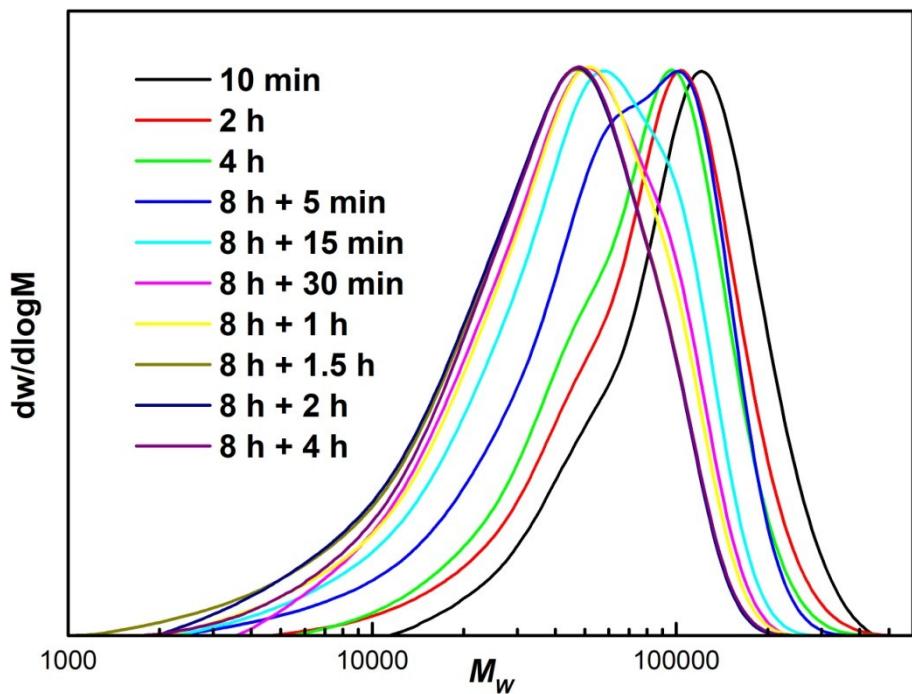


Figure 12S The GPC profiles evolution with the depolymerization (before 8 h) and cross metathesis time (after 8 h)

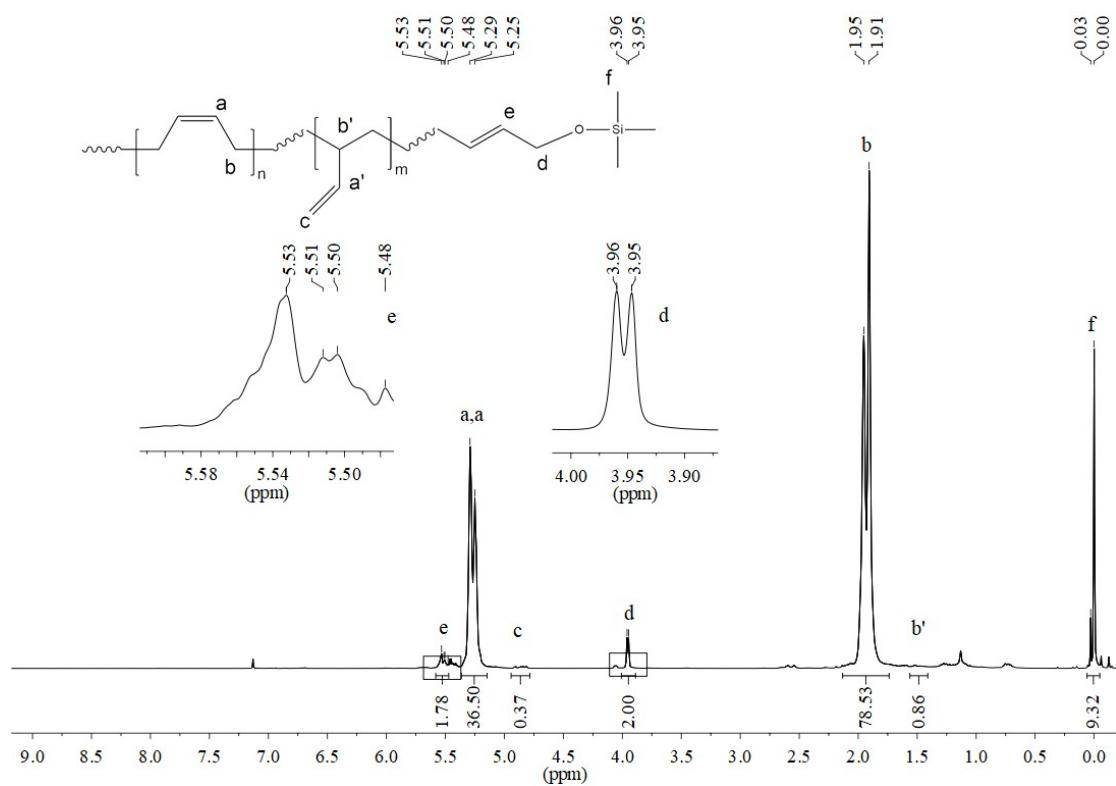


Figure 13S ^1H NMR spectra of PBD-SiMe₃.

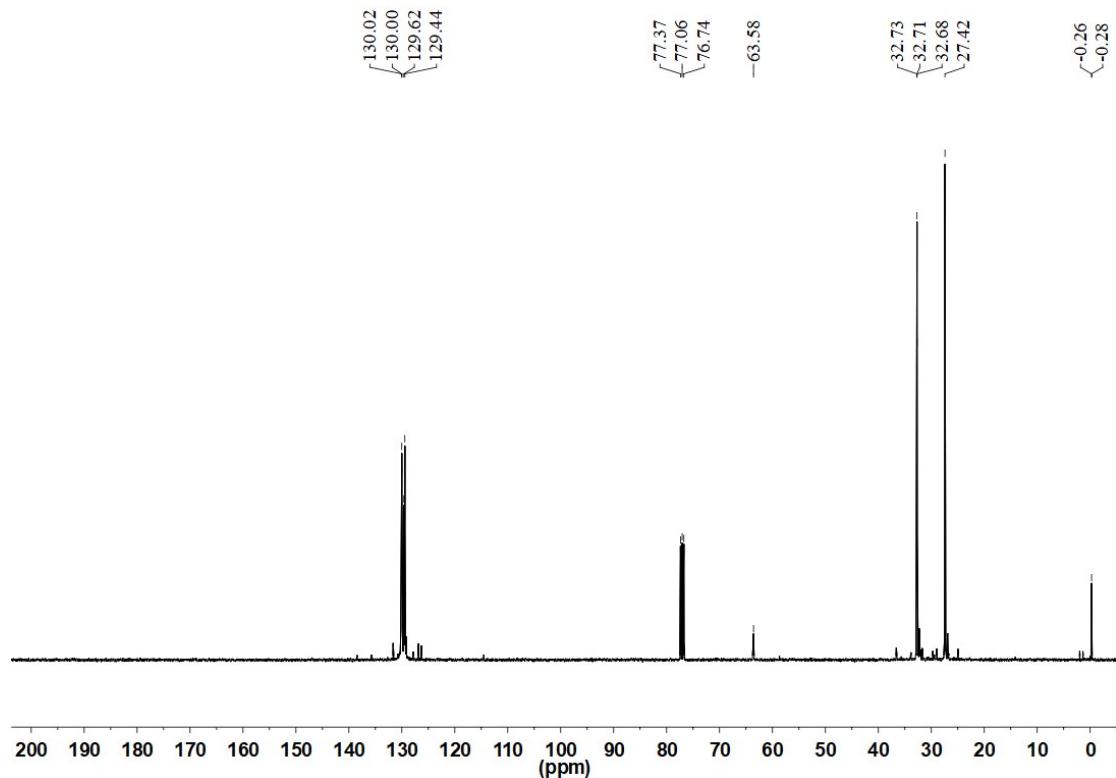


Figure 14S ^{13}C NMR spectra of PBD-SiMe₃.

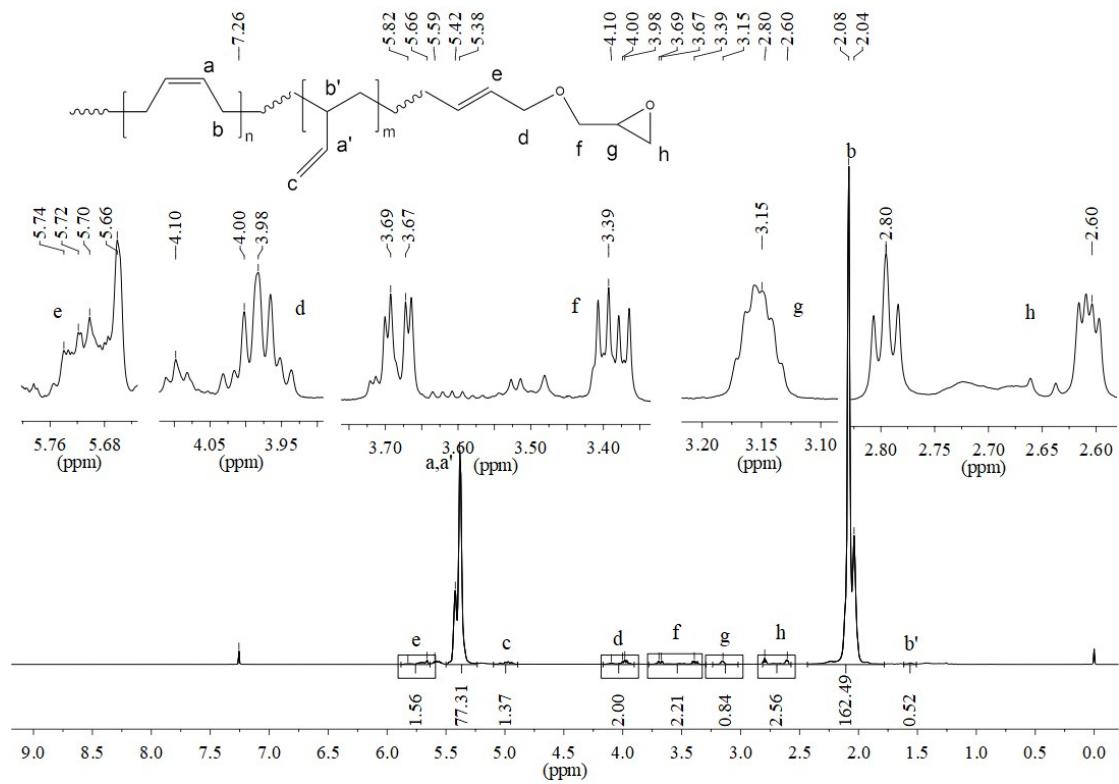


Figure 15S ^1H NMR spectra of PBD-PO

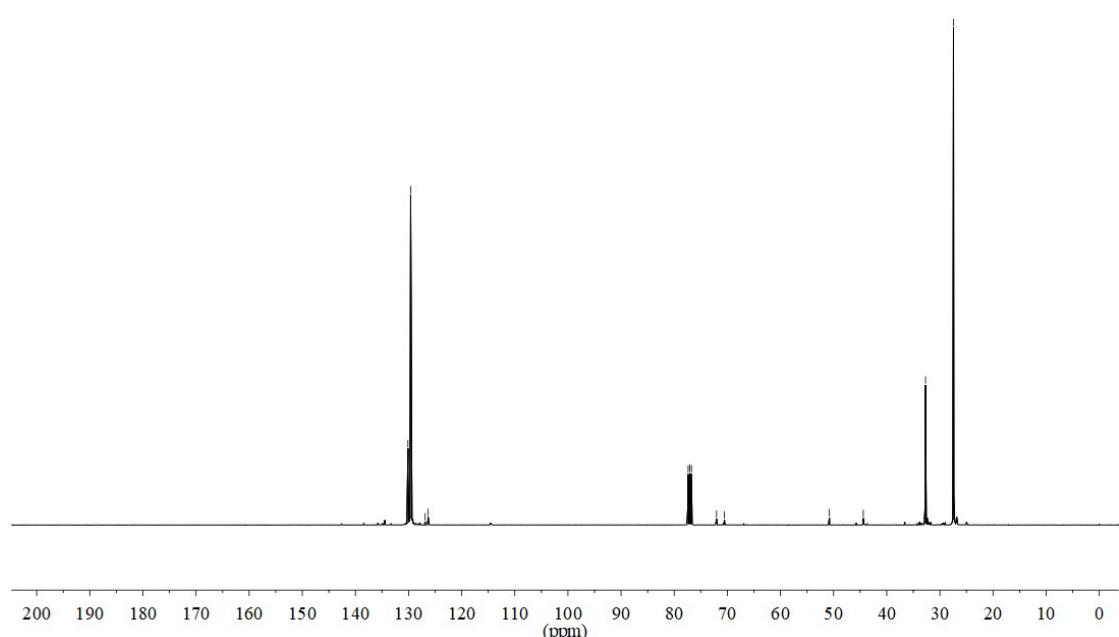


Figure 16S ^{13}C NMR spectra of PBD-PO

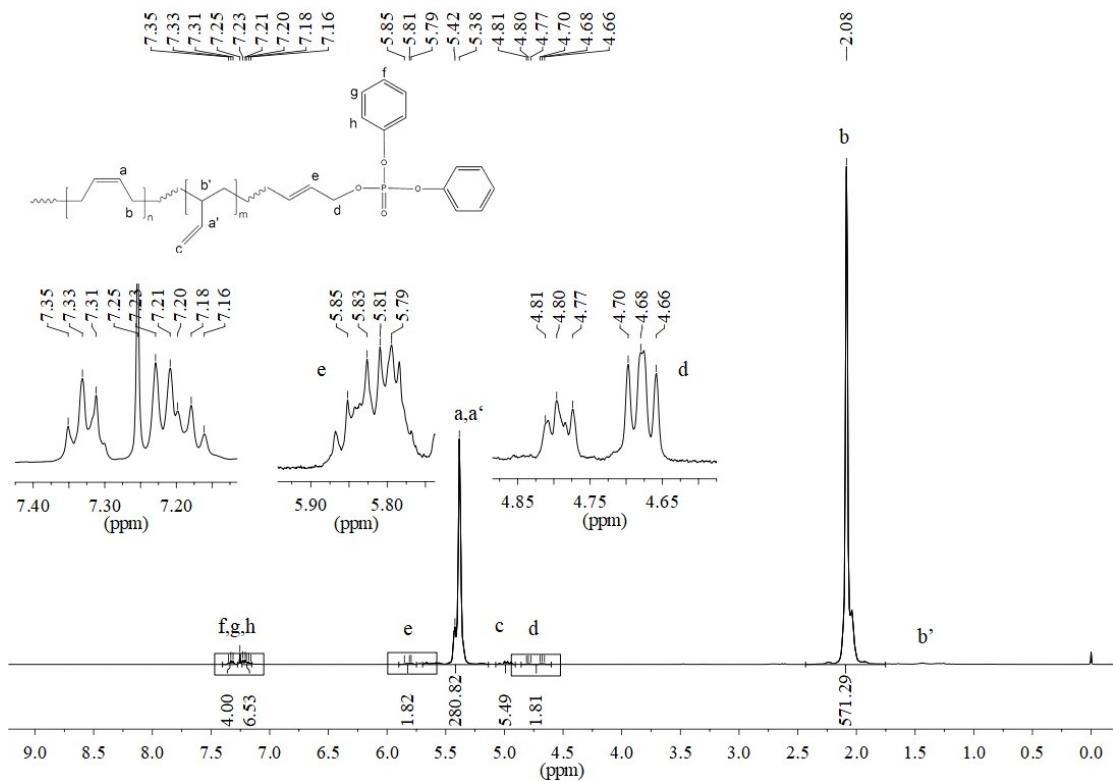


Figure 17S ^1H NMR spectra of PBD-Phosphate.

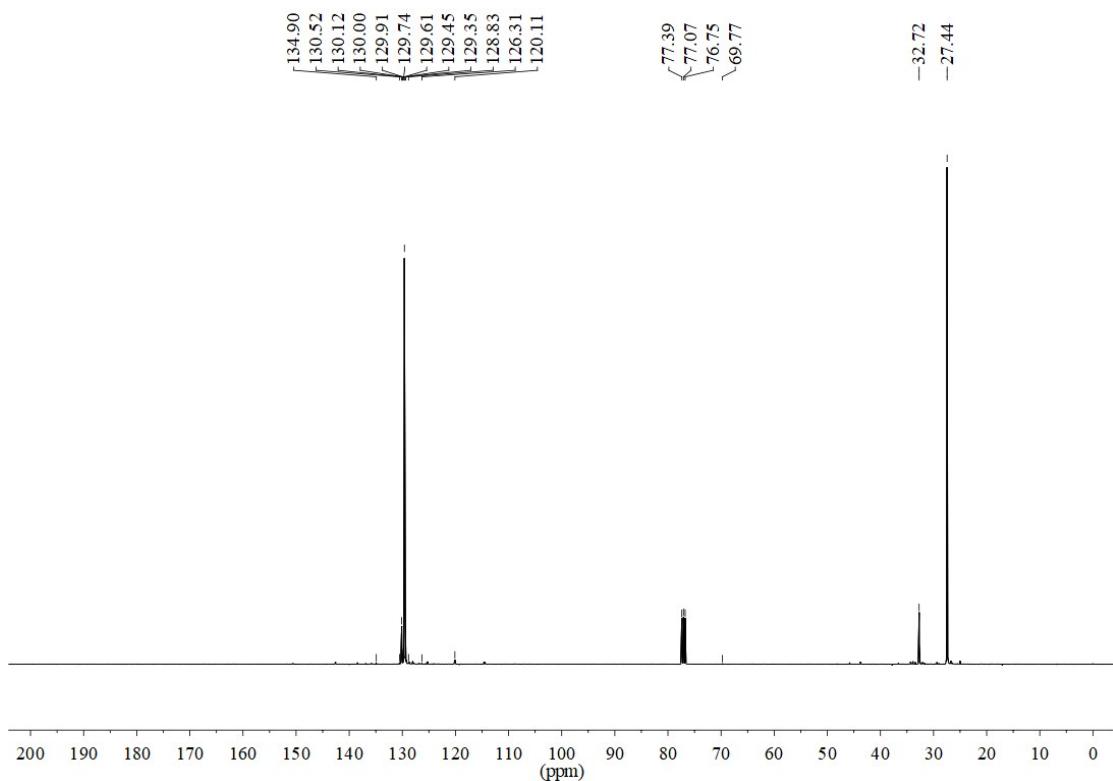


Figure 18S ^1H NMR spectra of PBD-Phosphate.

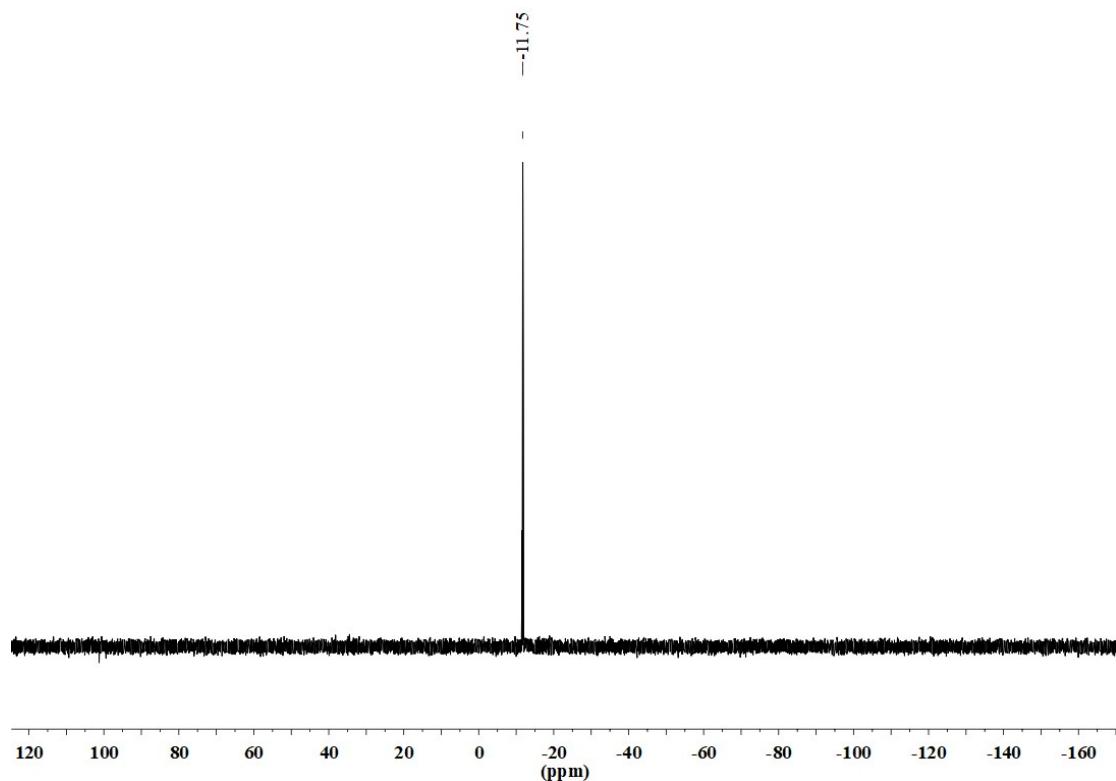


Figure 19S ^{31}P NMR spectra of PBD-Phosphate.

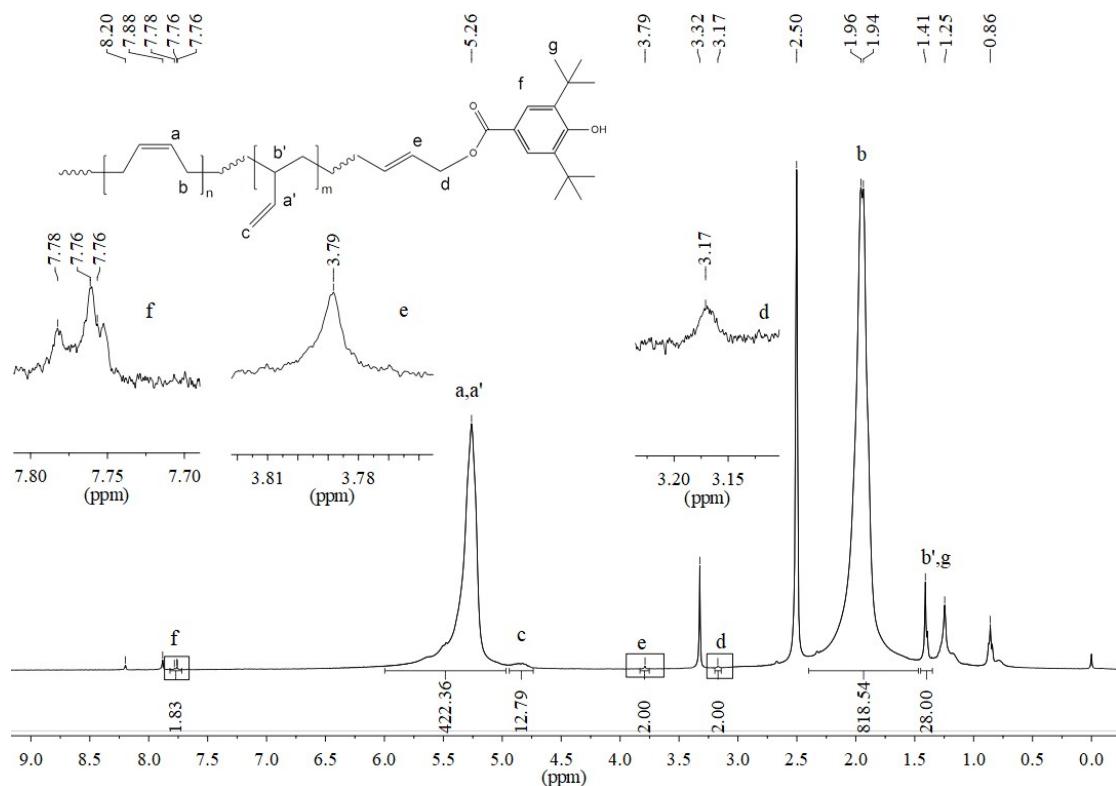


Figure 20S ^1H NMR spectra of PBD-Phenol

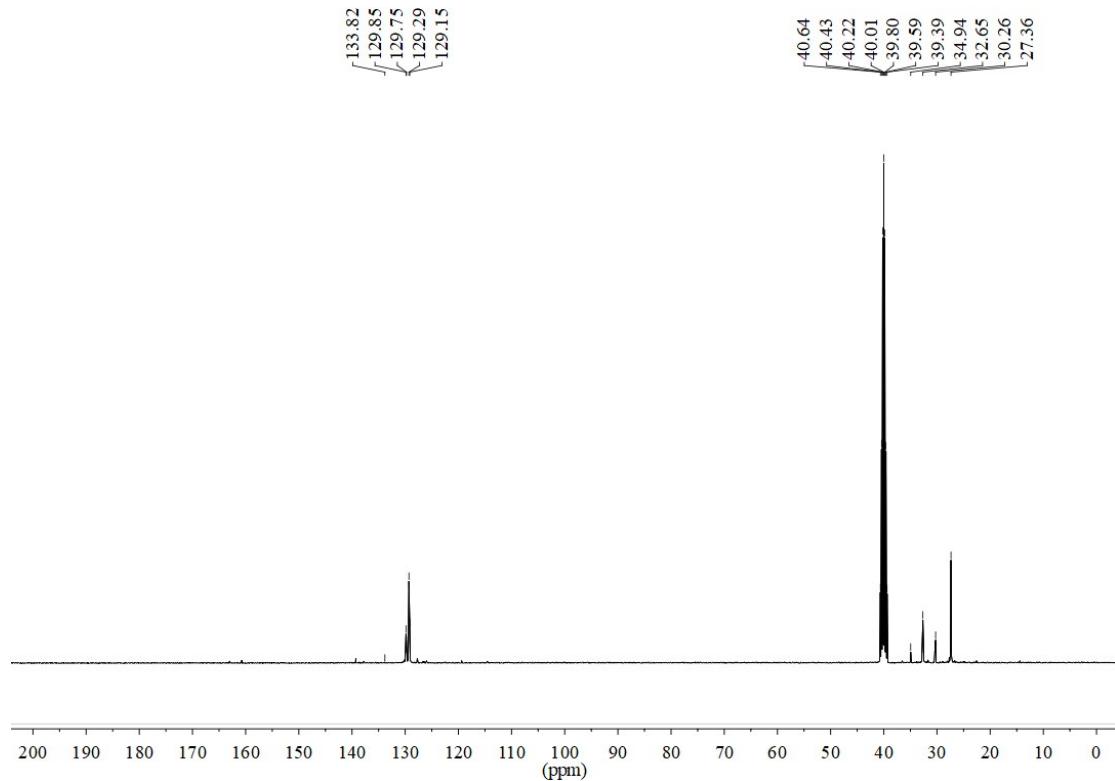


Figure 21S ¹³C NMR spectra of PBD-Phenol

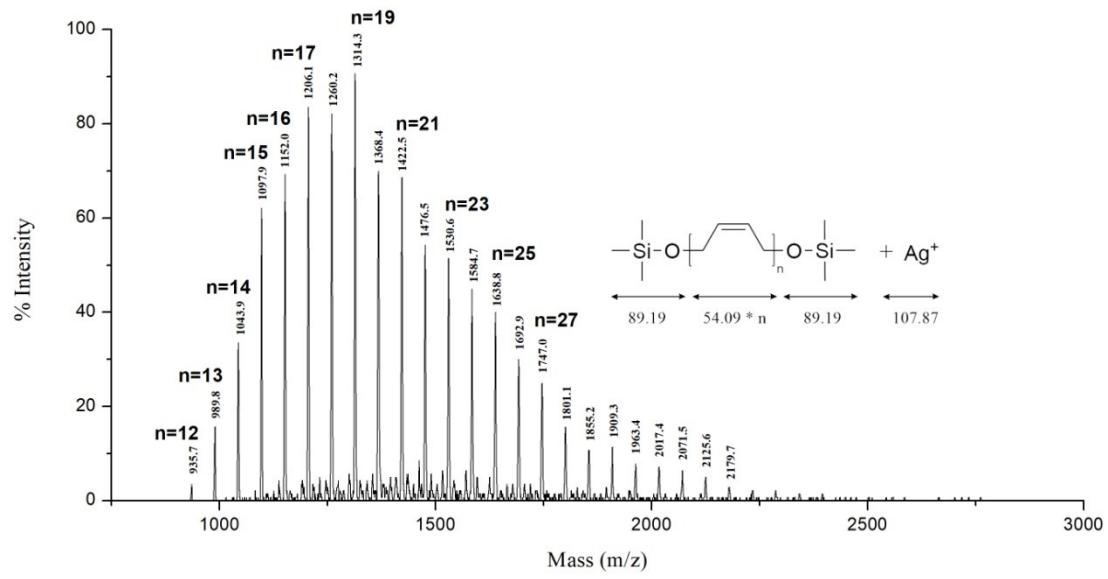


Figure 22S MALDI-ToF mass spectrum (DCTB matrix, silver ionizing salt) of PBD-SiMe₃

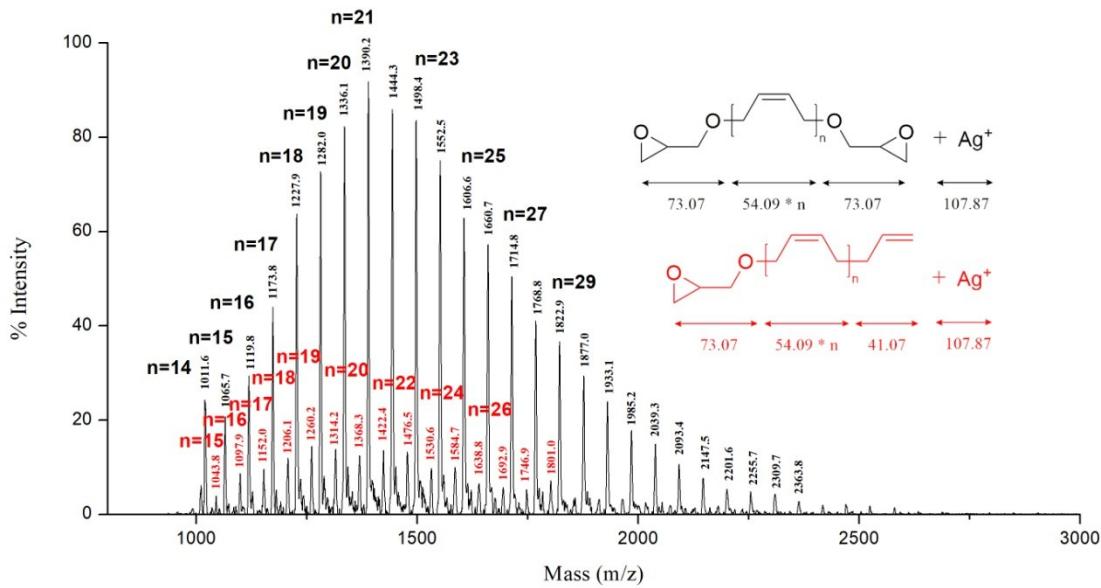


Figure 23S MALDI-ToF mass spectrum (DCTB matrix, silver ionizing salt) of **PBD-PO**

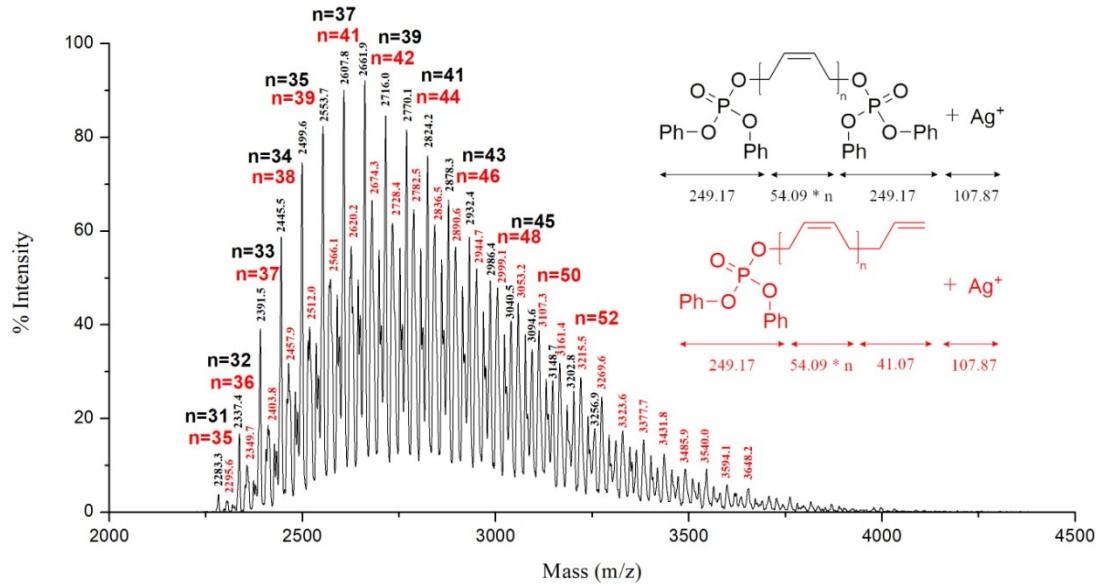


Figure 24S MALDI-ToF mass spectrum (DCTB matrix, silver ionizing salt) of **PBD-Phosphate**

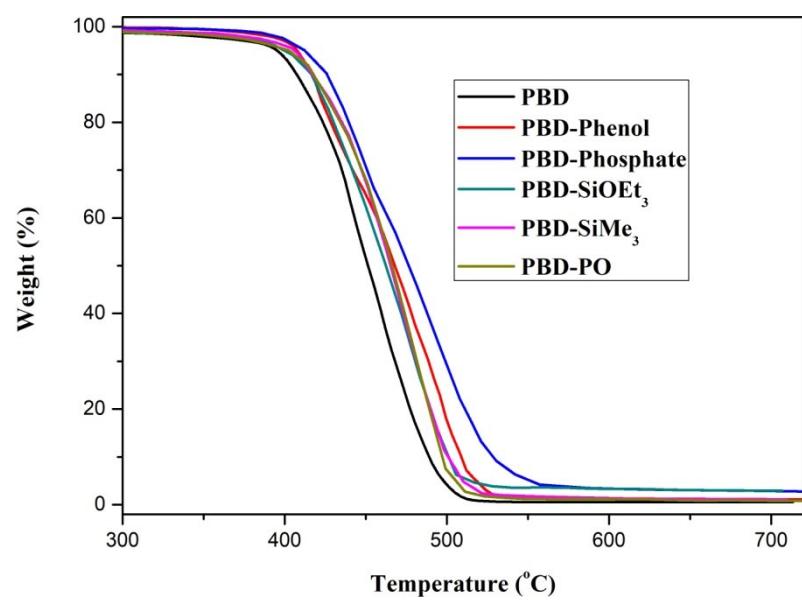


Figure 25S The TGA profiles of functionalized polybutadiene with various end groups

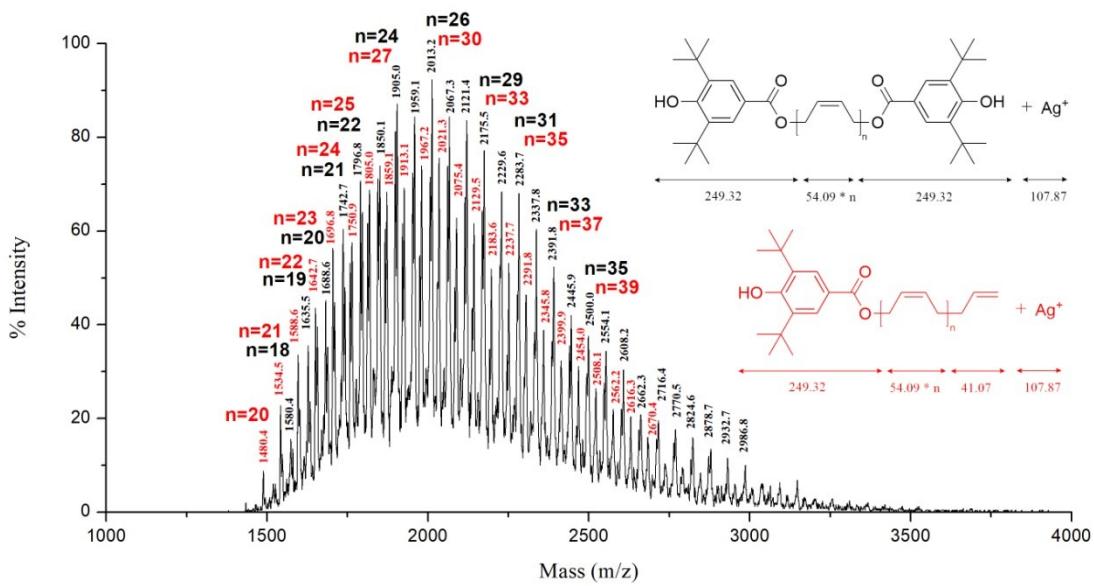


Figure 26S MALDI-ToF mass spectrum (DCTB matrix, silver ionizing salt) of **PBD-Phenol**

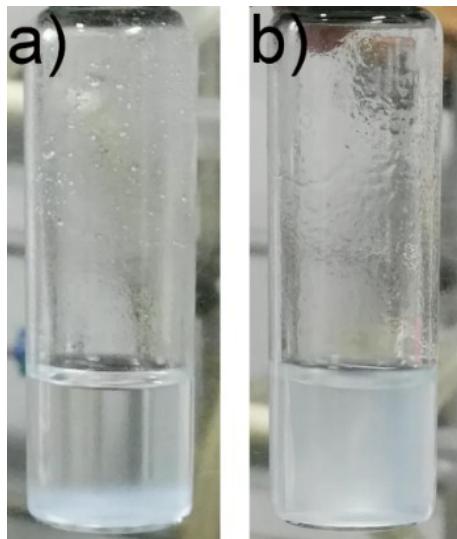


Figure 27S a) Si(OEt)₃-PBD modified SiO₂ suspension, after 24 h resting. (PBD-Si(OEt)₃: SiO₂ = 1:5), b) unmodified SiO₂ suspension in THF, after 24 h resting

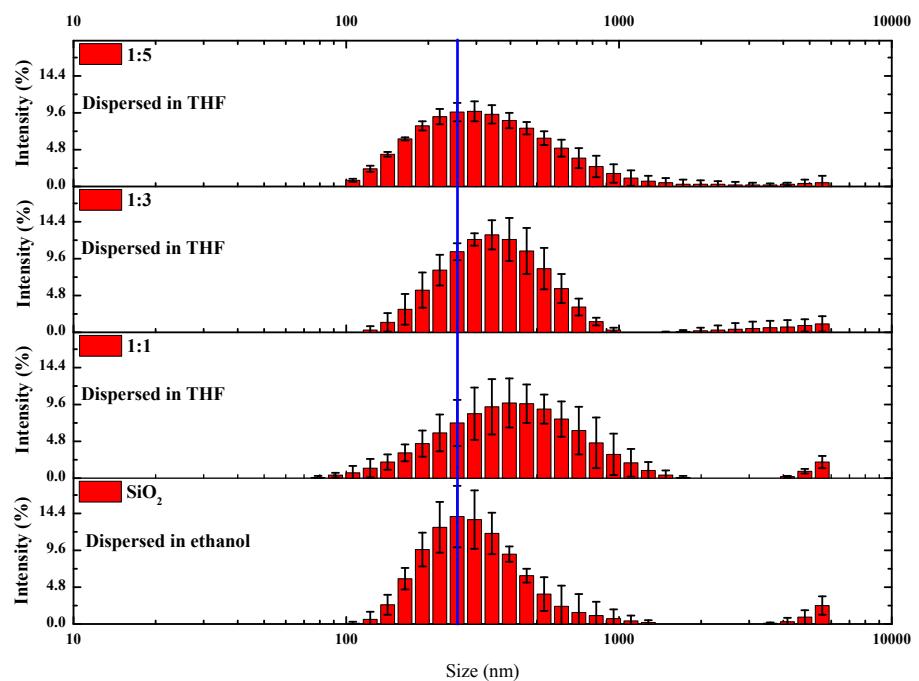


Figure 28S Particle size distribution of samples $\text{PBD-Si(OEt)}_3 : \text{SiO}_2 = 1:3, 1:5, 1:1$ in THF and pure SiO_2 in ethanol.

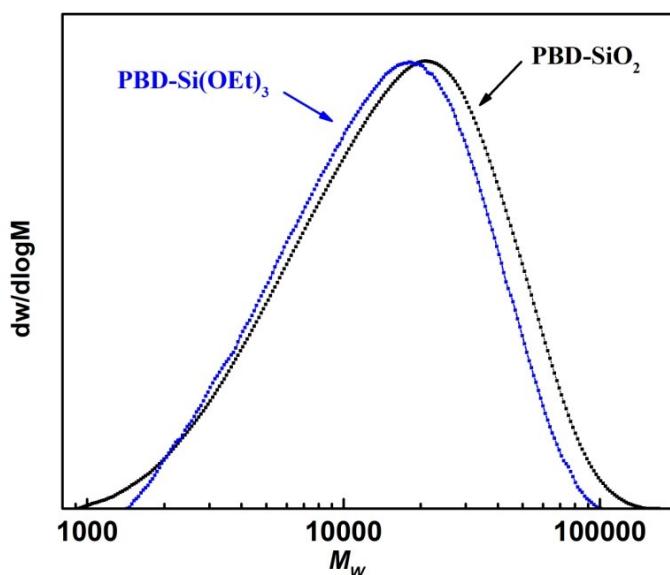


Figure 29S The GPC profile evolution of **PBD-Si(OEt)₃** (before hydrolysis on SiO_2) **PBD-SiO₂** (after hydrolysis on SiO_2).

Table 1S Composition of mixes.

	BR	10PSiO₂	10MPSiO₂	30PSiO₂	30MPSiO₂
Filler, phr	none	SiO_2 10	$\text{SiO}_2\text{-PBD}$ 10	SiO_2 30	$\text{SiO}_2\text{-PBD}$ 30
Base mix, phr	BR 100; ZnO 4;	Stearic acid 2;	Sulphur 1.5;	CZ 1.5, D 0.5;	4010NA 1

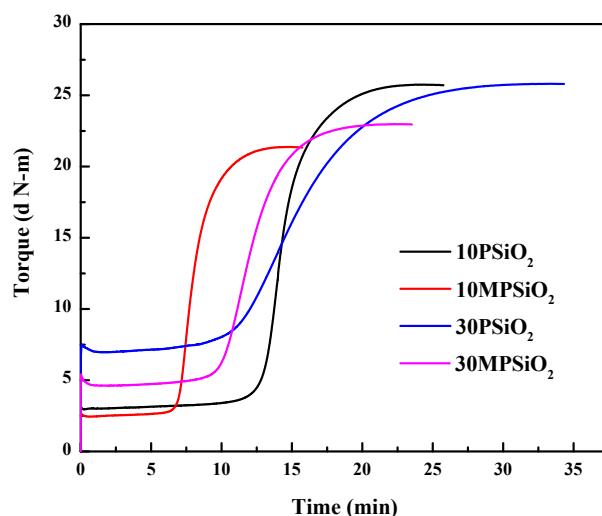


Figure 30S Cure rheographs of PBD blends containing pure SiO_2 and $\text{SiO}_2\text{-PBD}$ at 150 $^{\circ}\text{C}$