

Self-Healing UV-Curable Urethane (Meth)acrylates with Various Soft Segment Chemistry

Paulina Bednarczyk ^{1,*}, Paula Ossowicz-Rupniewska ¹, Joanna Klebeko ¹, Joanna Rokicka ¹, Yongping Bai ^{2,3} and Zbigniew Czech ¹

¹ Department of Chemical Organic Technology and Polymeric Materials, Faculty of Chemical Technology and Engineering, West Pomeranian University of Technology in Szczecin, Piastów Ave. 42, 71-065 Szczecin, Poland; possowicz@zut.edu.pl (P.O.-R.); joanna.klebeko@zut.edu.pl (J.K.); psa_czech@wp.pl (Z.C.)

² School of Chemistry and Chemical Engineering, Harbin Institute of Technology, Harbin 150001, China

³ Wuxi HIT New Material Research Institute Co., Ltd., Wuxi 214000, China

* Correspondence: bednarczyk.pb@gmail.com or paulina.bednarczyk@zut.edu.pl

HODA (Hydroxymethyl)-10-oxatricyclo[5.2.1.0^{2,6}]dec-8-ene-3,5-dione-2-aminoethanol

¹H NMR (400 MHz, CDCl₃, 298 K) δ (ppm) 6.61 (d, 1H), 6.55 (t, 1H), 5.28 (s, 1H), 4.11 (s, 2H), 3.76 (t, 2H), 3.69 (t, 2H), 3.04 (d, 1H), 3.01 (d, 1H), 2.85 (s, 1H), 2.28 (s, 1H); ¹³C NMR (400 MHz, CDCl₃, 298 K) δ (ppm): 138.31; 136.96; 91.50; 80.96; 60.72; 60.16; 49.99; 48.17; 41.75; FT-IR (cm⁻¹) 417, 456, 499, 522, 584, 613, 644, 678, 716, 731, 787, 805, 836, 844, 876, 932, 953, 977, 1012, 1036, 1074, 1091, 1111, 1159, 1185, 1248, 1272, 1289, 1303, 1326, 1348, 1363, 1402, 1428, 1681, 1766, 2883, 2917, 2929, 2950, 2978, 3016, 3085, 3428.

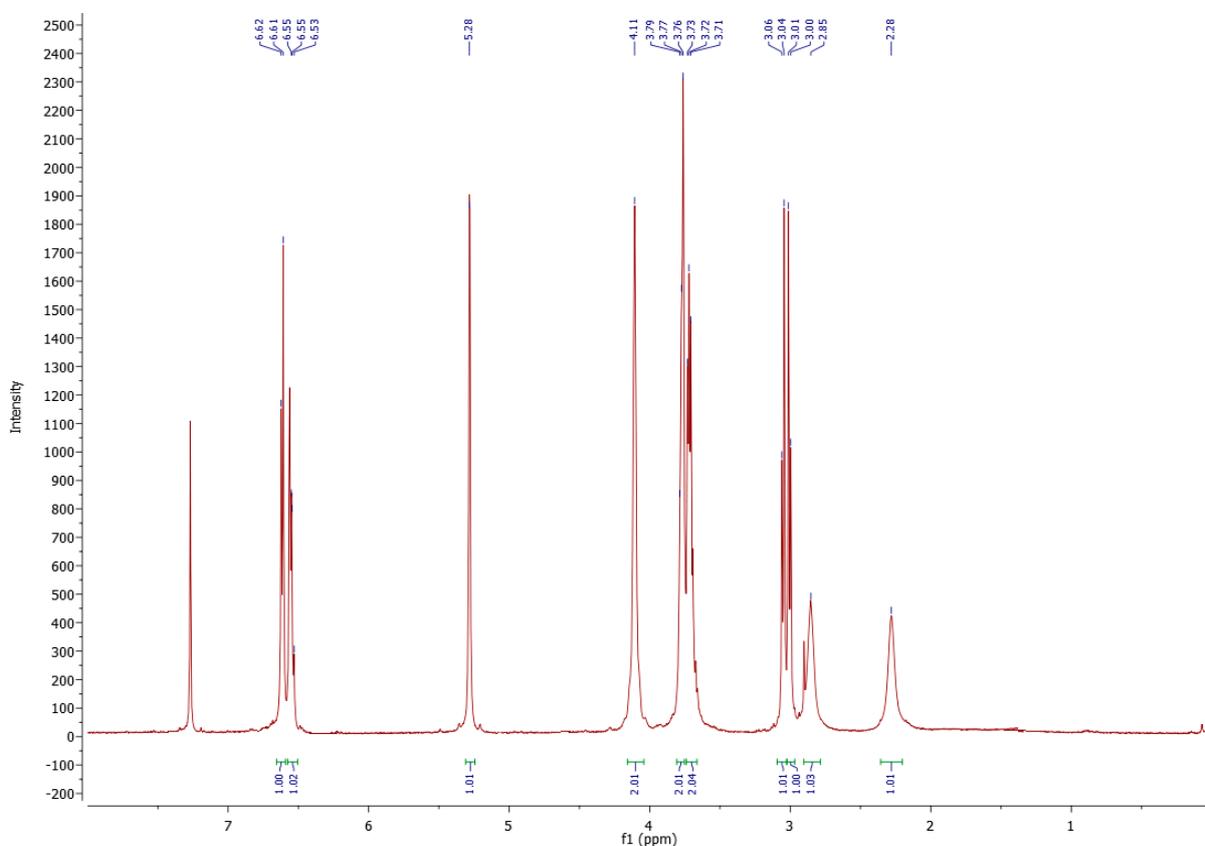


Figure S1. ¹H NMR HODA spectrum.

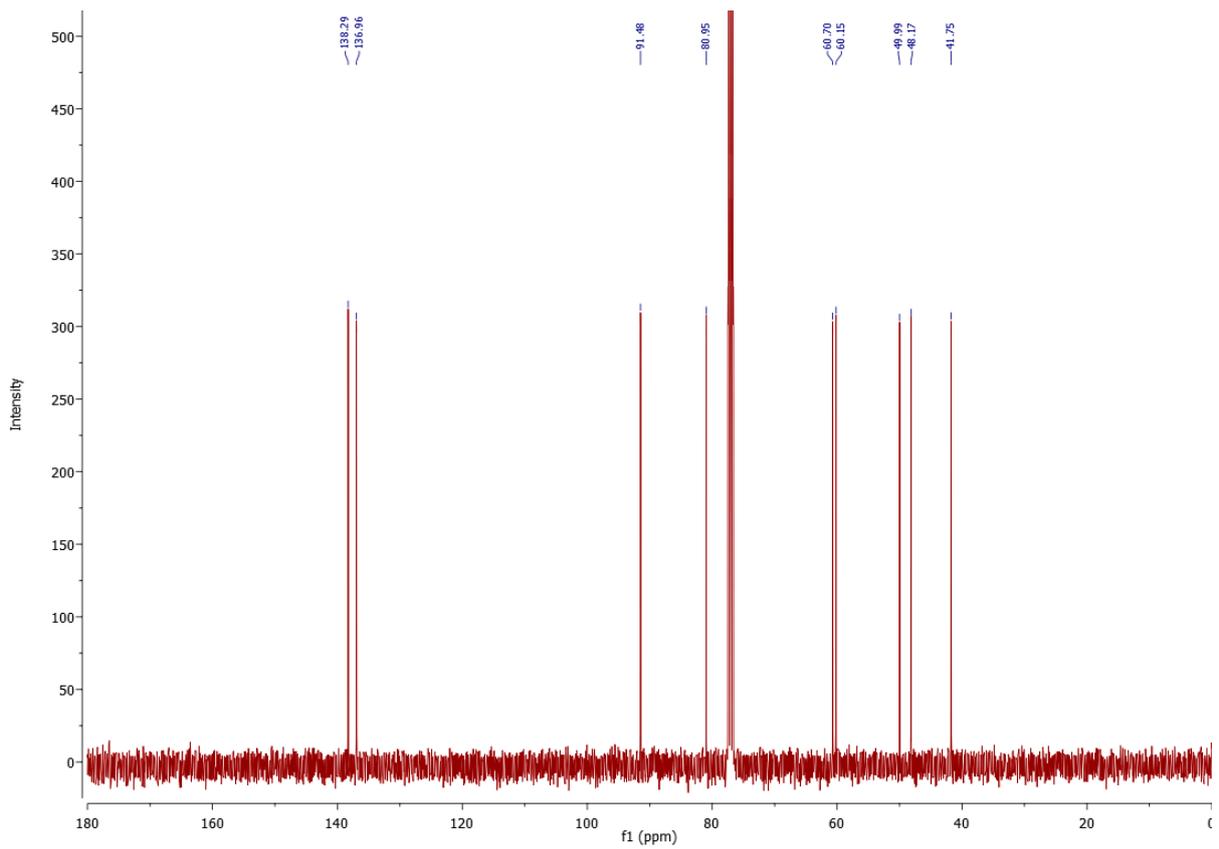


Figure S2. ¹³C NMR HODA spectrum.

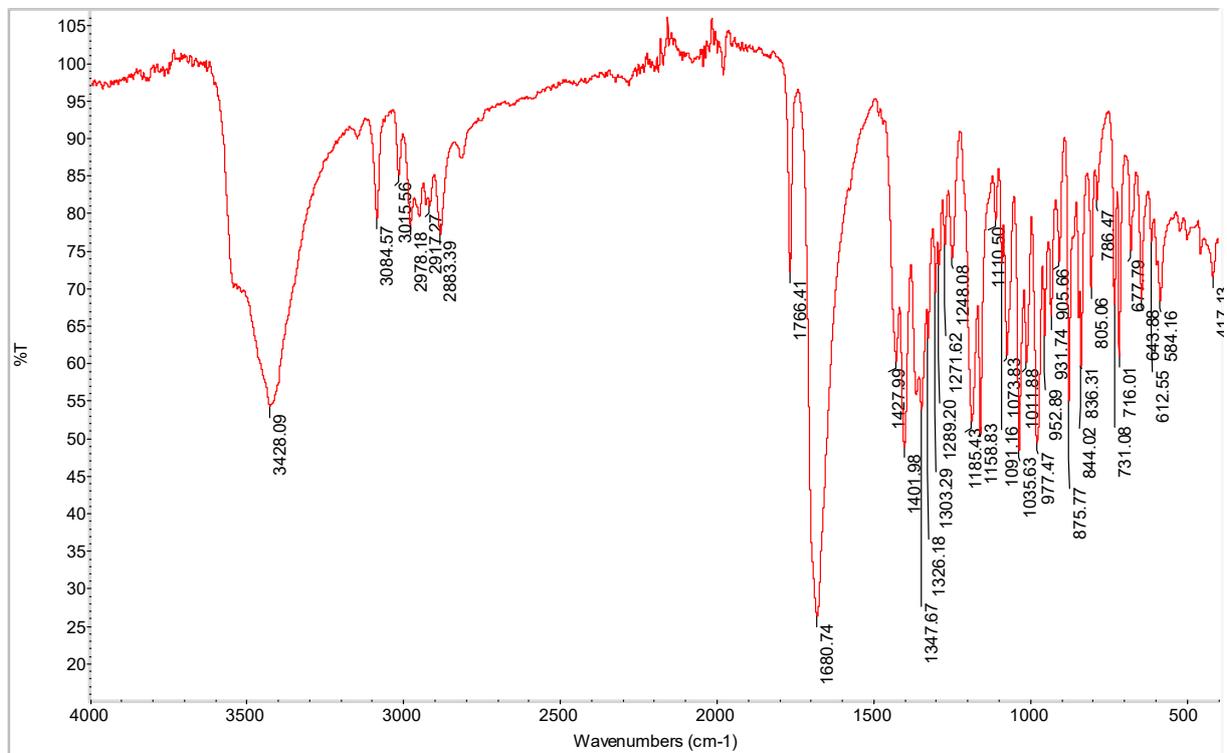


Figure S3. ATR-FTIR HODA spectrum.

**The FT-IR spectra during the reaction time for the obtained urethane
(meth)acrylates**

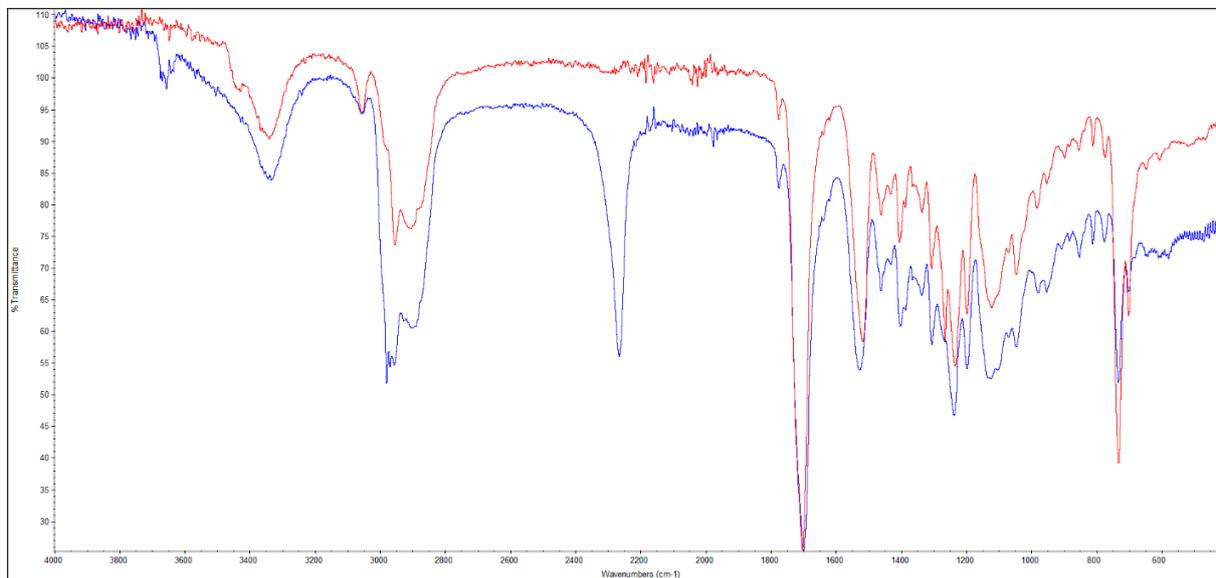


Figure S4. Comparative of FTIR spectra investigating the reaction of receiving PEG400-HPA after 0 (blue) and 1 h (red) of reaction time. Comparison of FTIR peaks: NCO stretching at 2270 cm^{-1} .

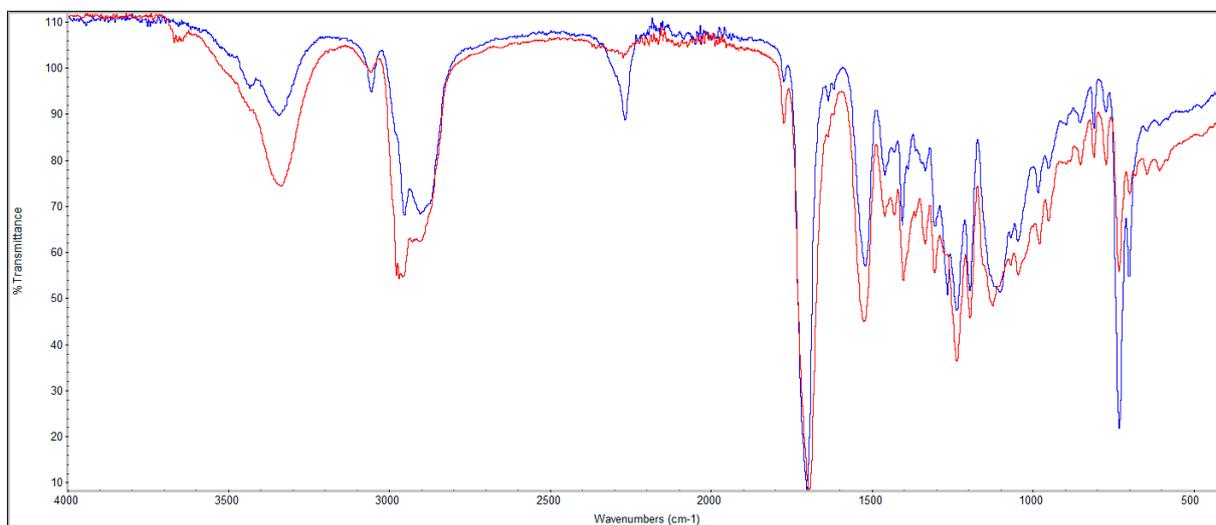


Figure S5. Comparative of FTIR spectra investigating the reaction of receiving PEG600-HPA after 0 (blue) and 1 h (red) of reaction time. Comparison of FTIR peaks: NCO stretching at 2270 cm^{-1} .

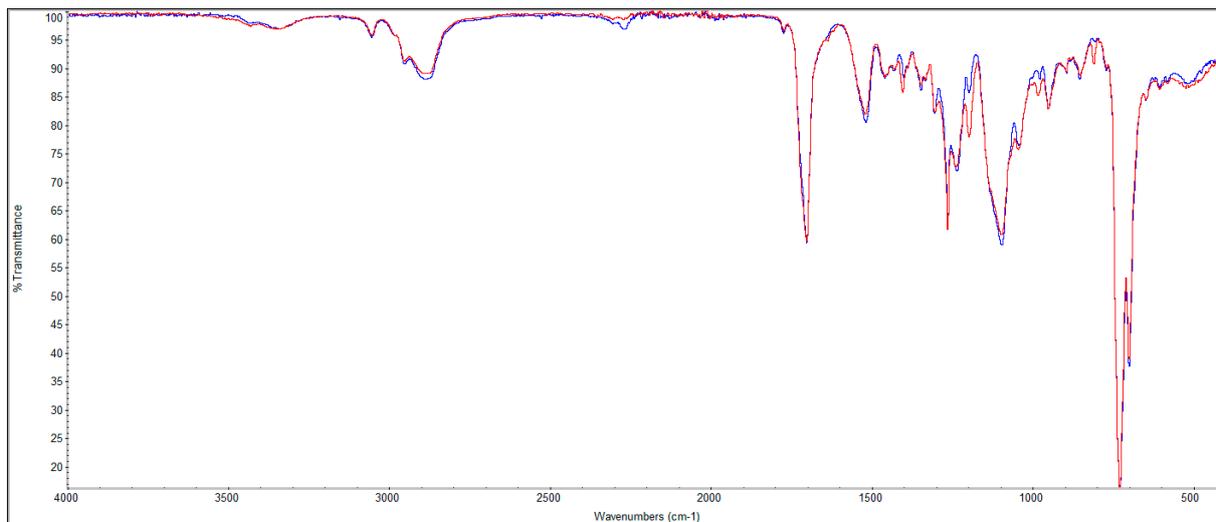


Figure S6. Comparative of FTIR spectra investigating the reaction of receiving PEG1000-HPA after 0 (blue) and 1 h (red) of reaction time. Comparison of FTIR peaks: NCO stretching at 2270 cm⁻¹.

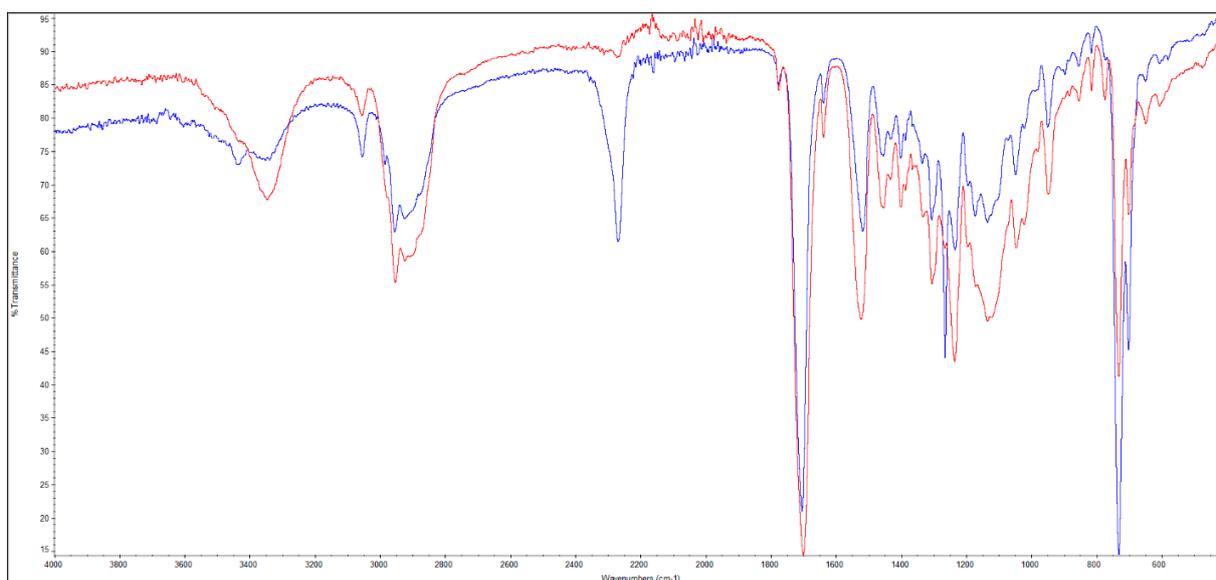


Figure S7. Comparison of FTIR spectra investigating the reaction of receiving PEG400-HPMA after 0 (blue) and 3 h (red) reaction time. Comparison of FTIR peaks: NCO stretching at 2270 cm⁻¹.

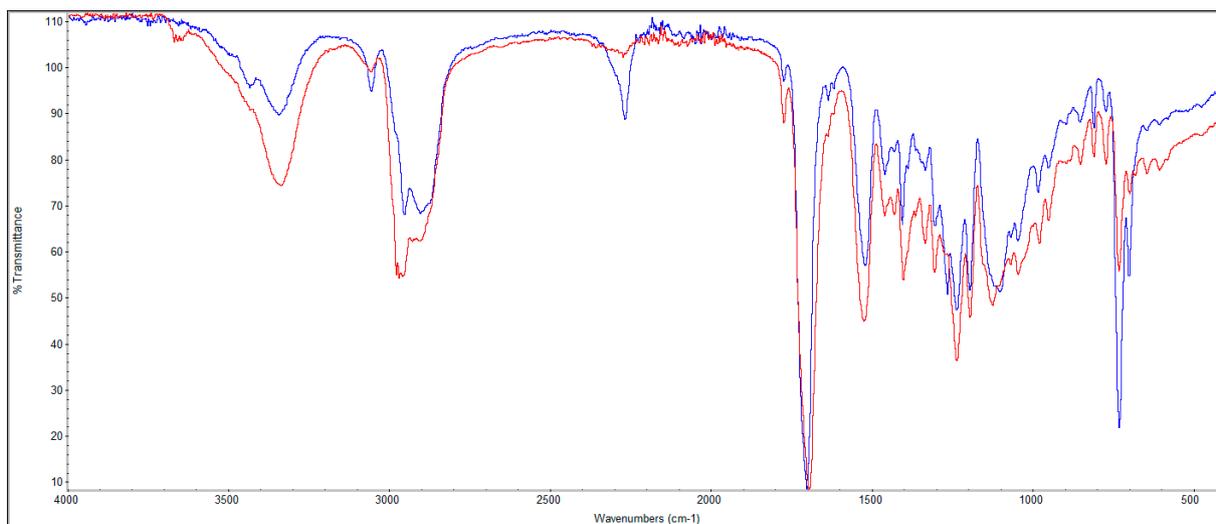


Figure S8. Comparative of FTIR spectra investigating the reaction of receiving PEG600-HPMA after 0 (blue) and 4 h (red) of reaction time. Comparison of FTIR peaks: NCO stretching at 2270 cm^{-1} .

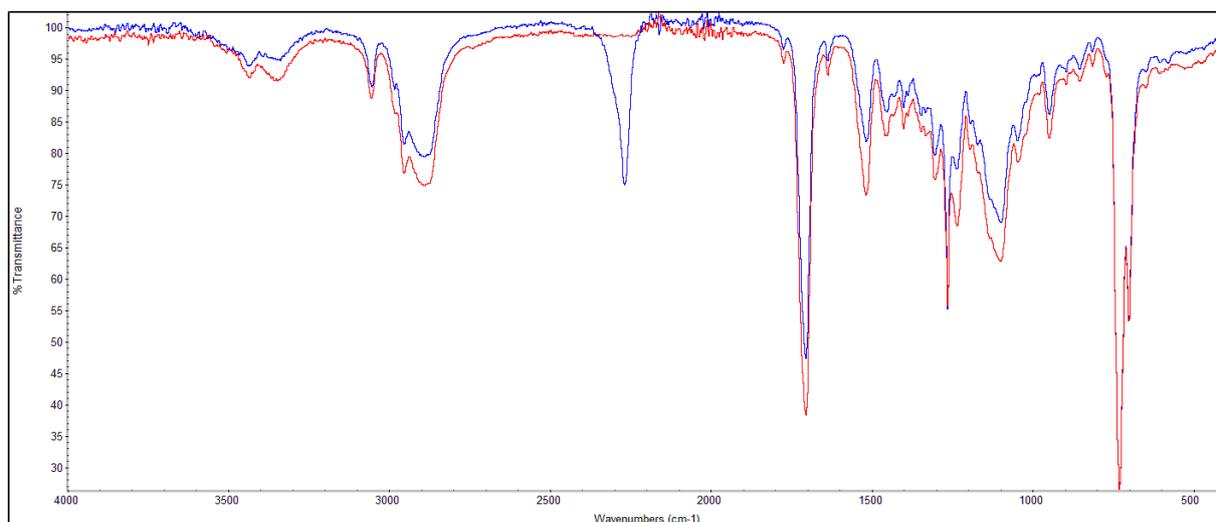


Figure S9. Comparative of FTIR spectra investigating the reaction of receiving PEG1000-HPMA after 0 (blue) and 3 h (red) of reaction time. Comparison of FTIR peaks: NCO stretching at 2270 cm^{-1} .

Selected NMR spectra of the obtained urethane (meth)acrylates

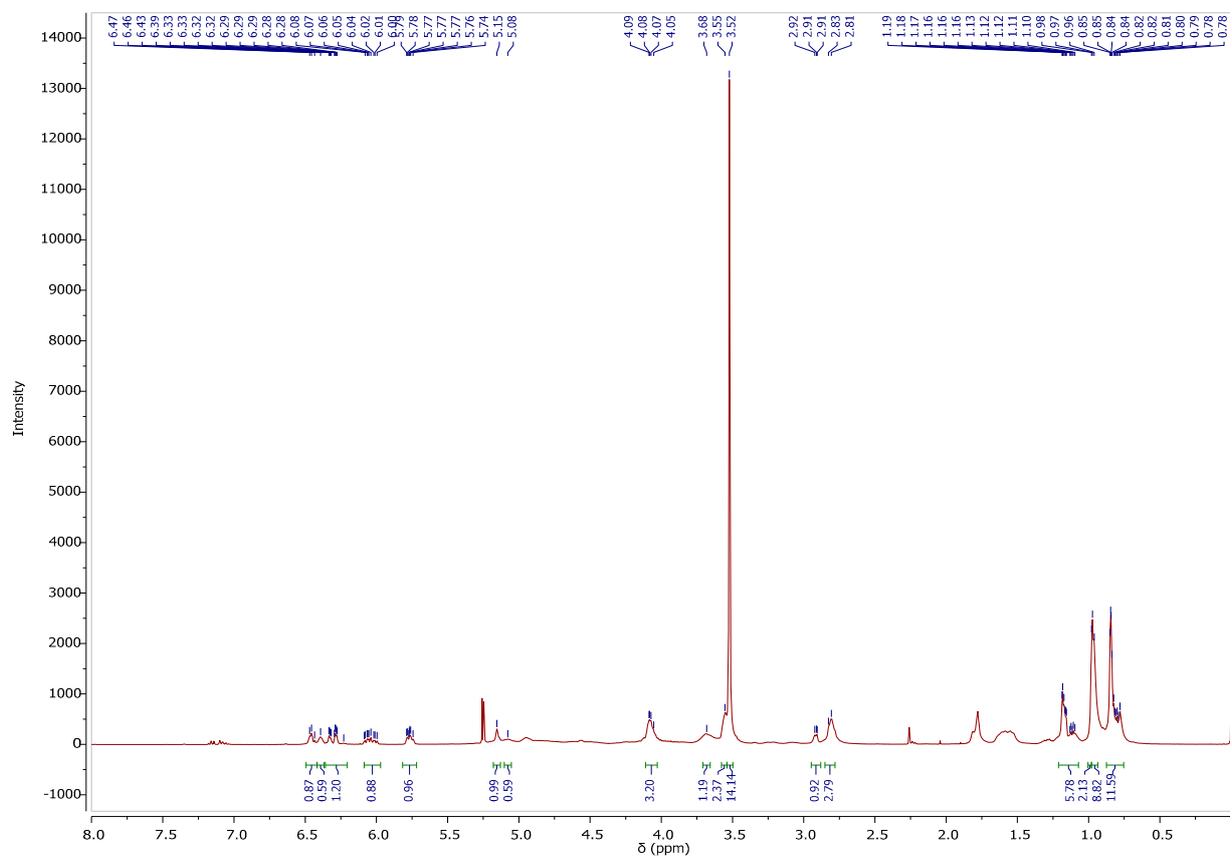


Figure S10. ¹H NMR spectra of PEG400-HPA.

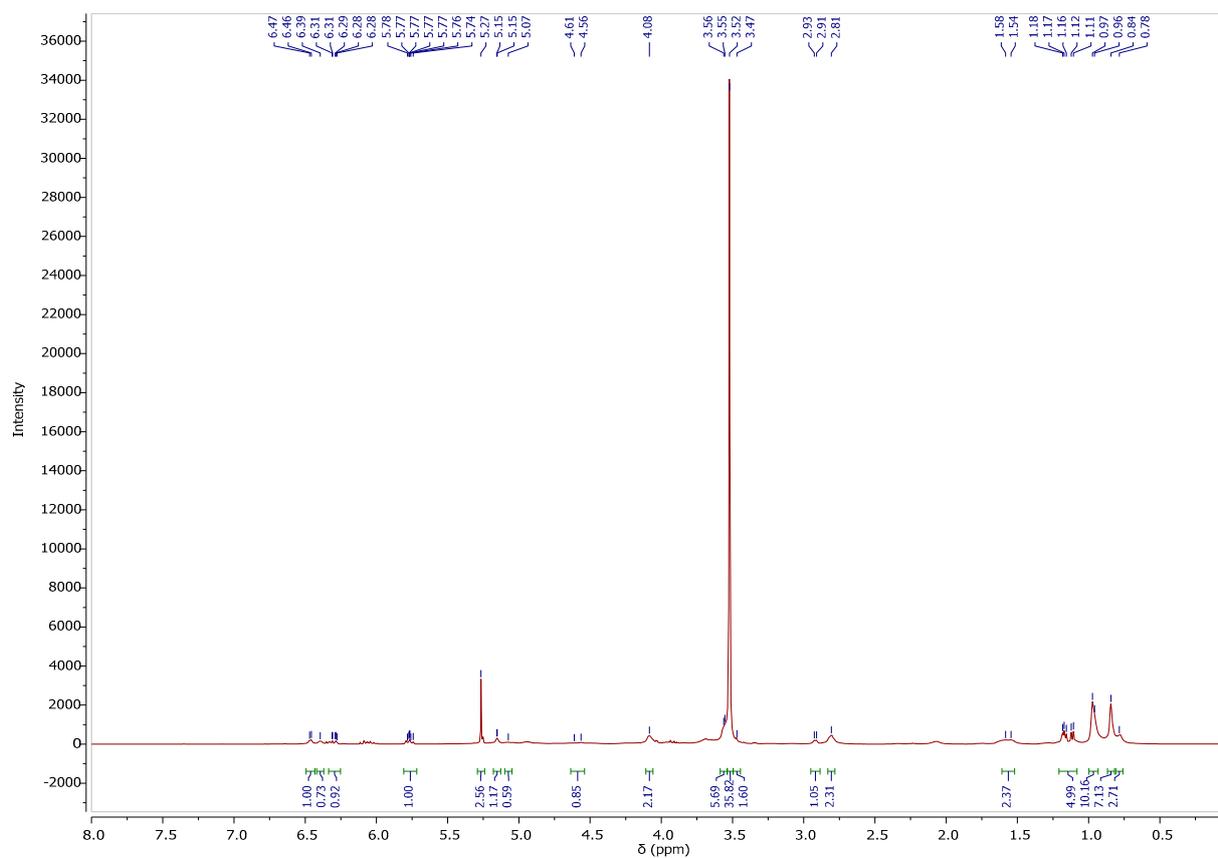


Figure S12. ^1H NMR spectra of PEG1000-HPA.

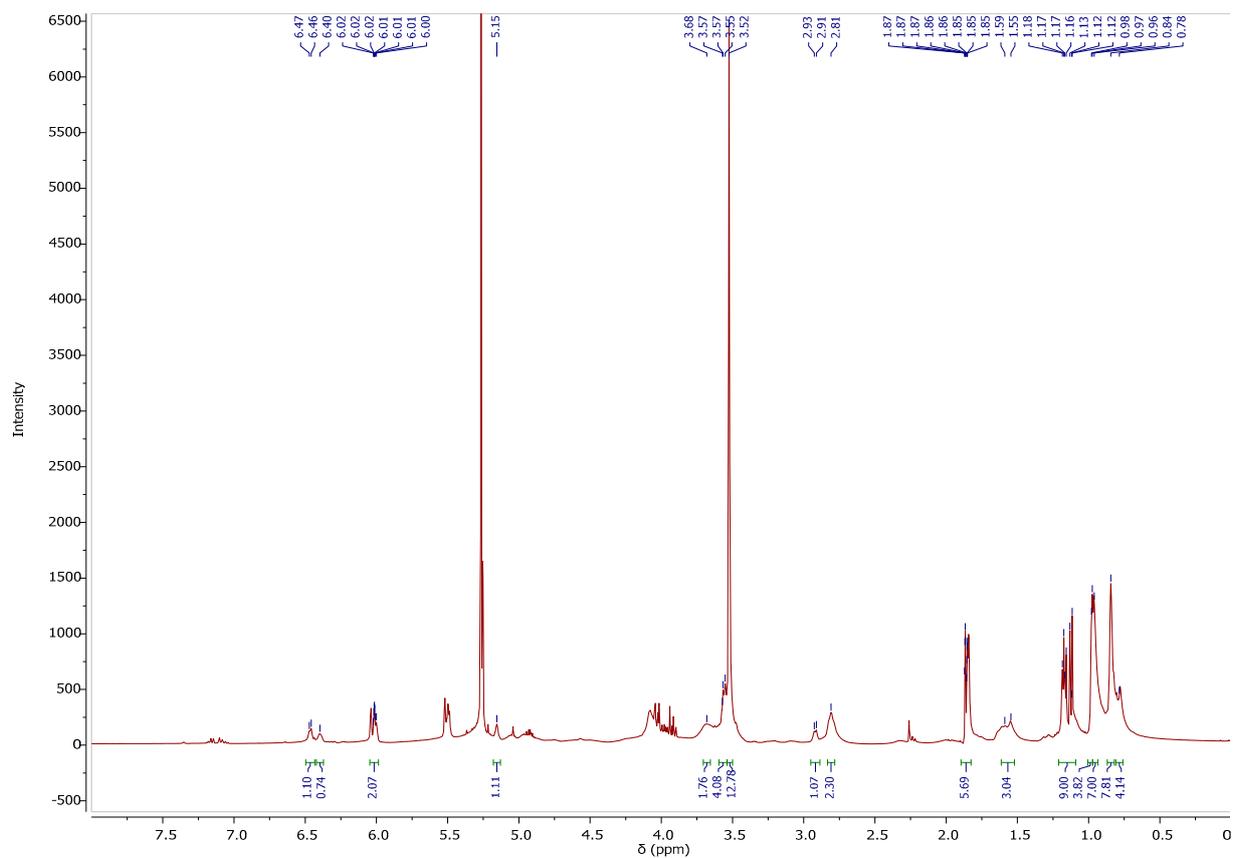


Figure S13. ^1H NMR spectra of PEG400-HPMA.

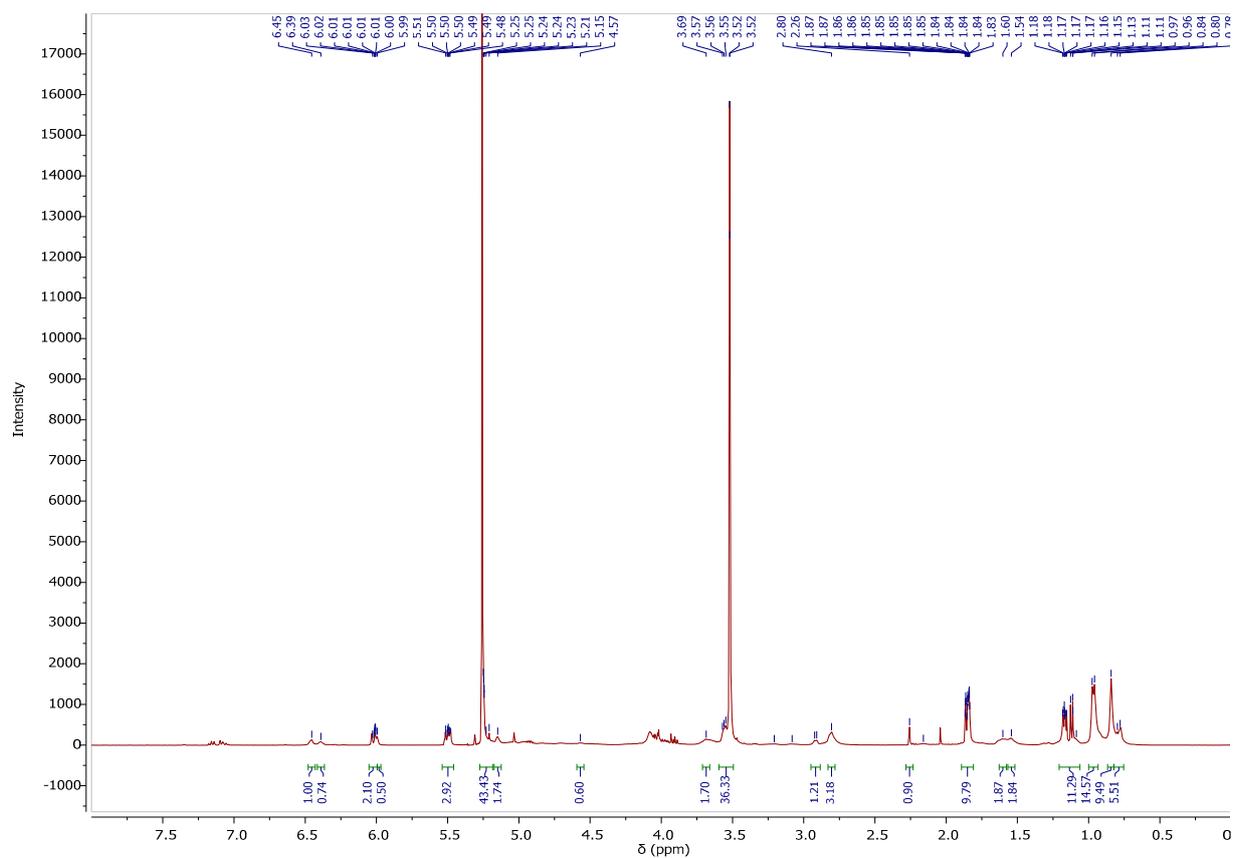


Figure S14. ¹H NMR spectra of PEG600-HPMA.

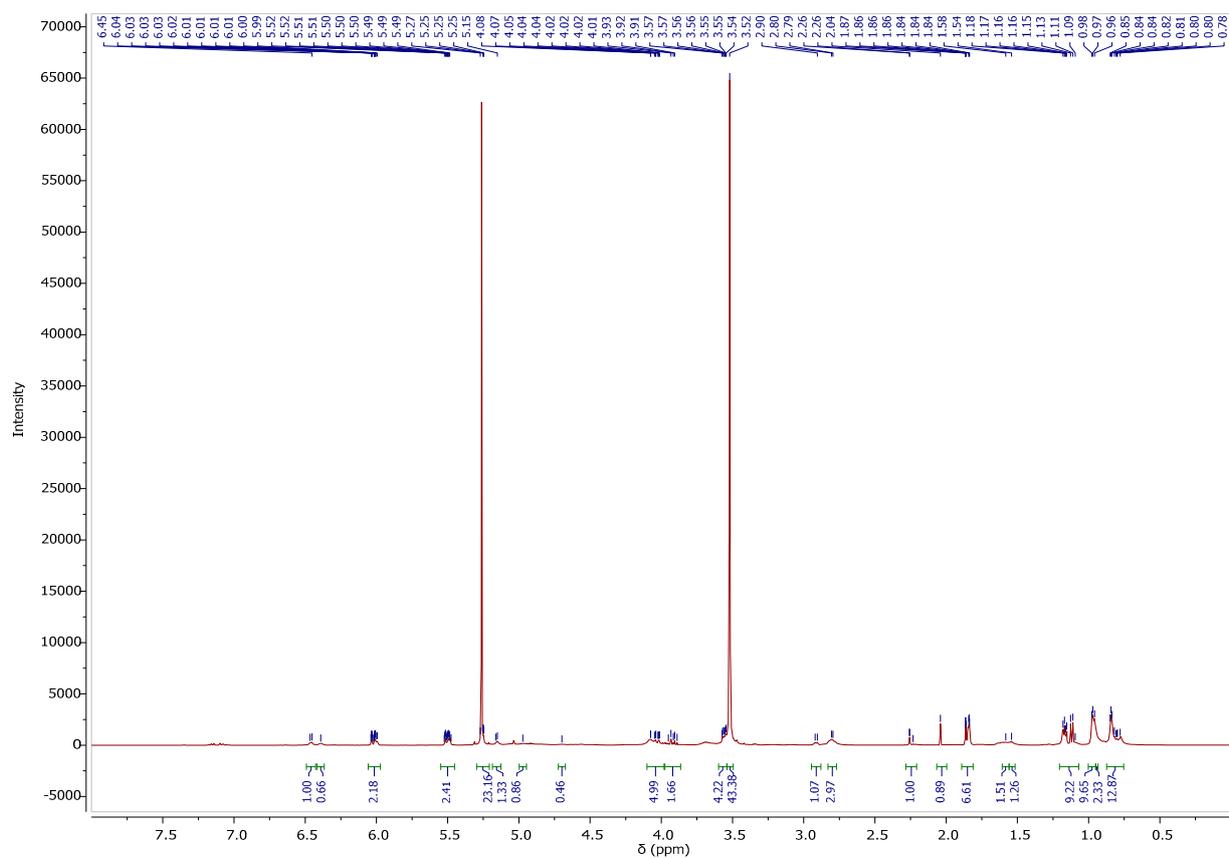


Figure S15. ^1H NMR spectra of PEG1000-HPMA.

The TG and DTG curves of the obtained urethane (meth)acrylates.

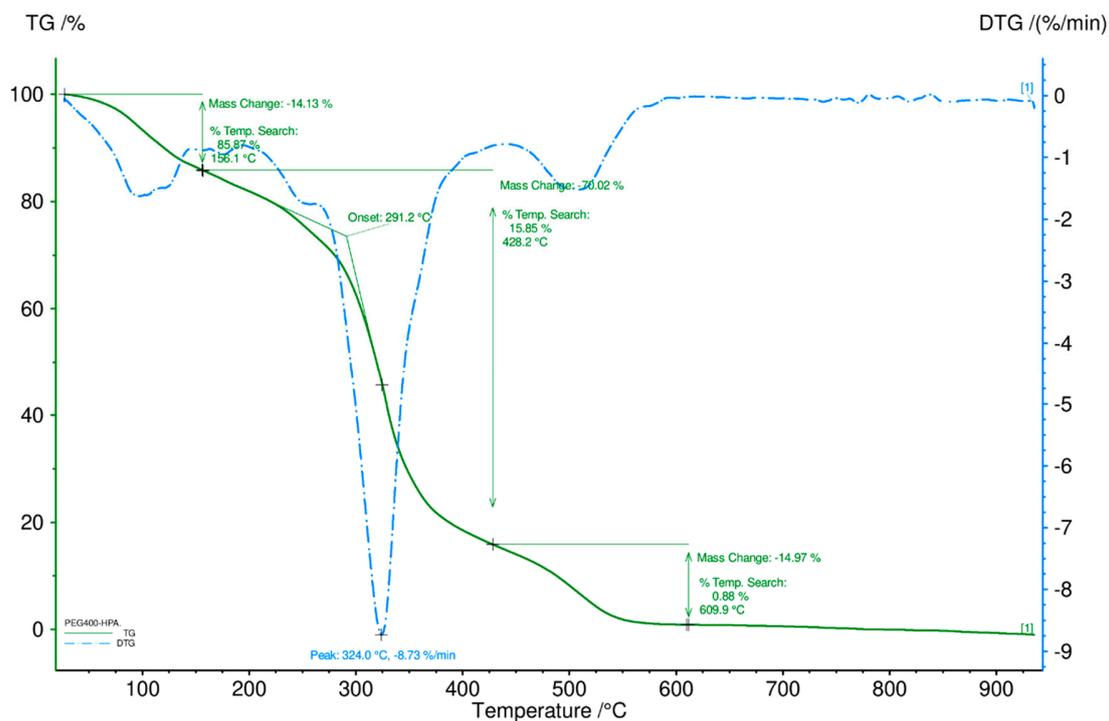


Figure S16. TG and DTG curves of PEG400-HPA.

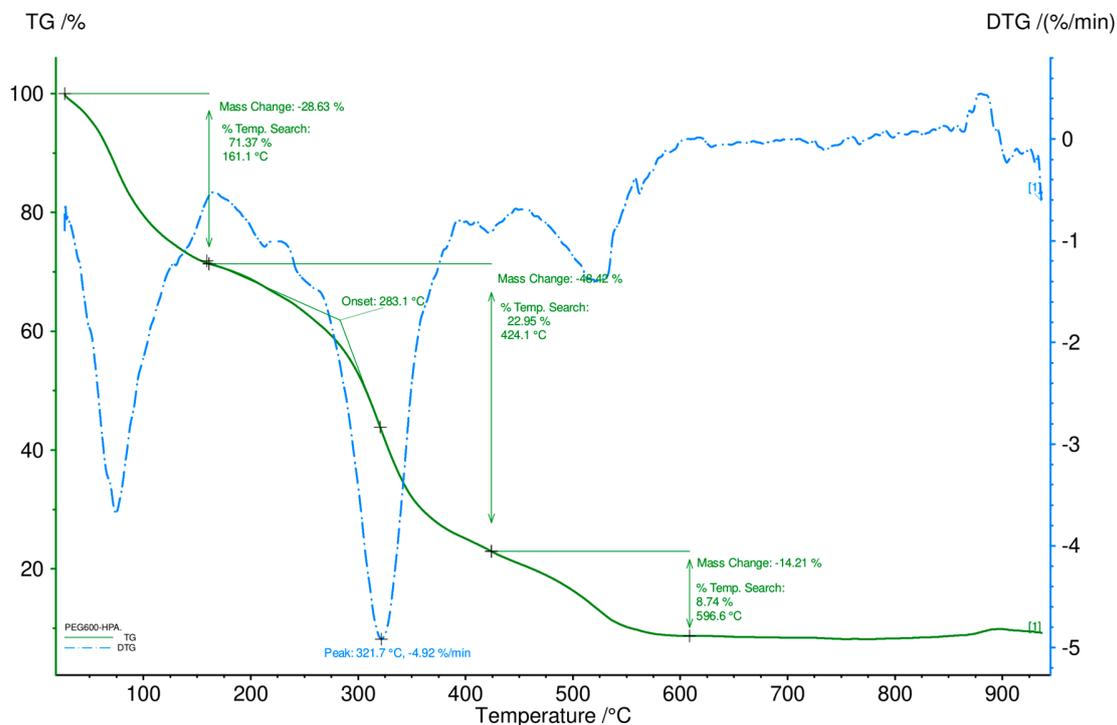


Figure S17. TG and DTG curves of PEG600-HPA.

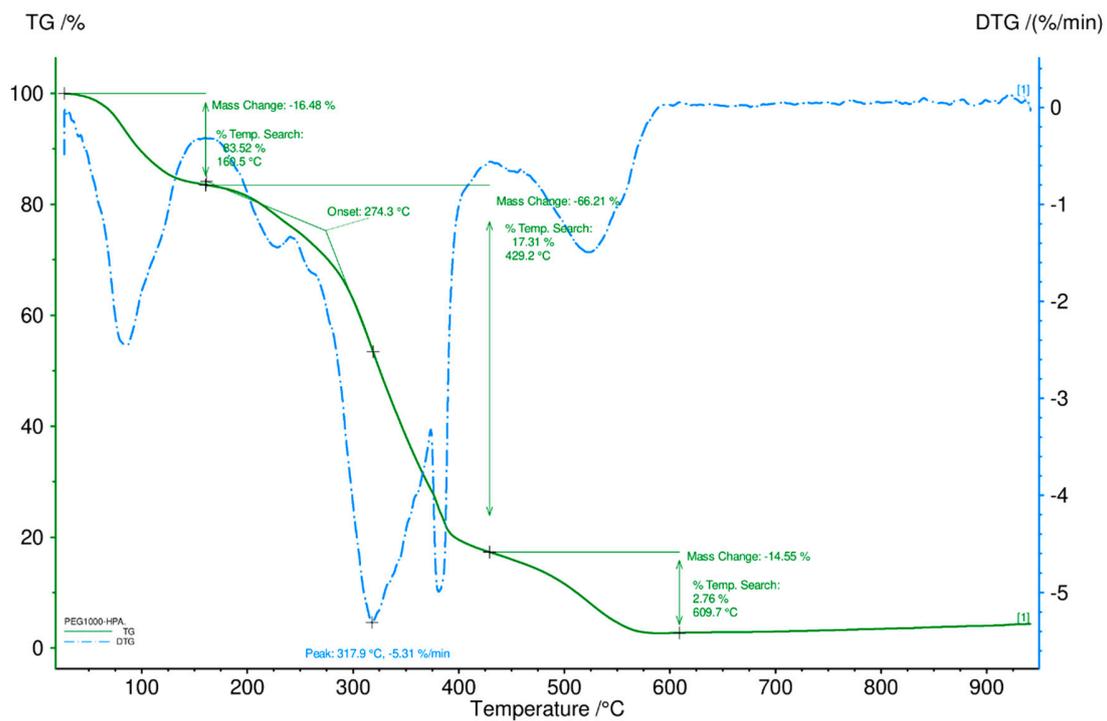


Figure S17. TG and DTG curves of PEG1000-HPA.

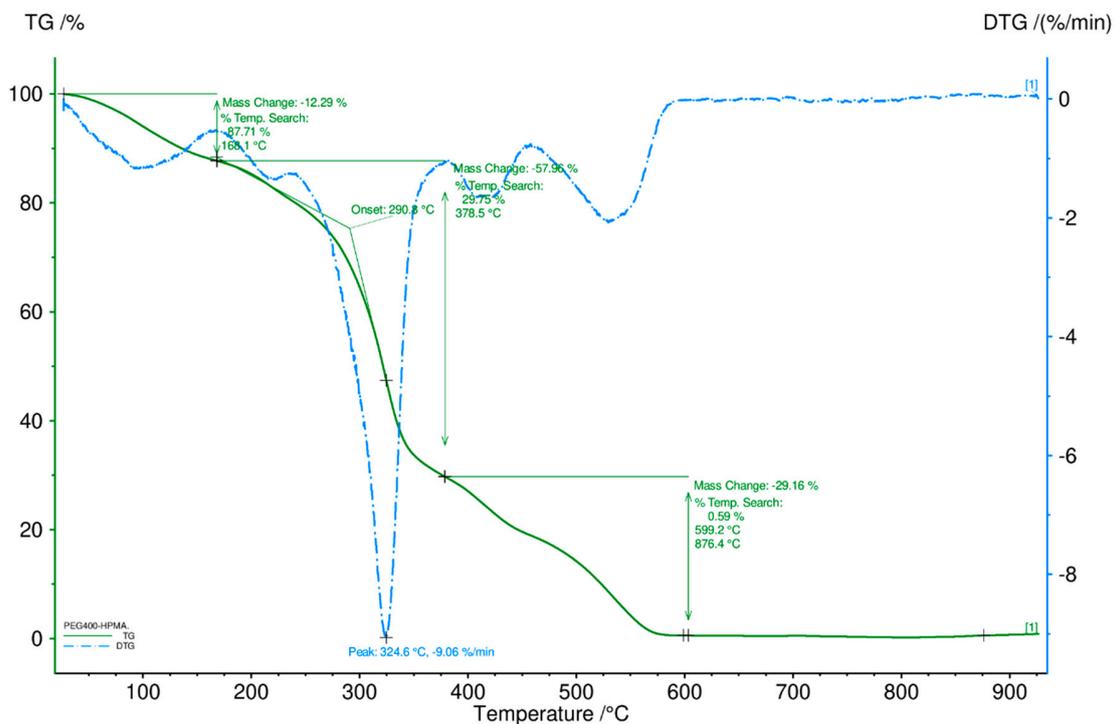


Figure S19. TG and DTG curves of PEG400-HPMA.

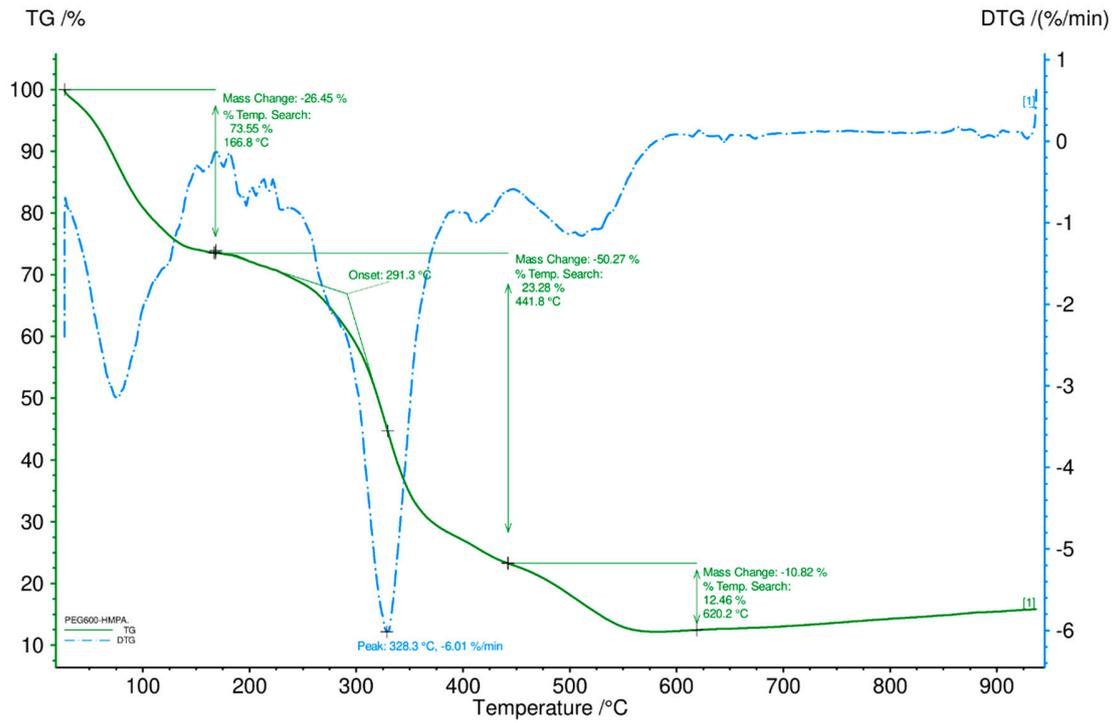


Figure S18. TG and DTG curves of PEG600-HPMA.

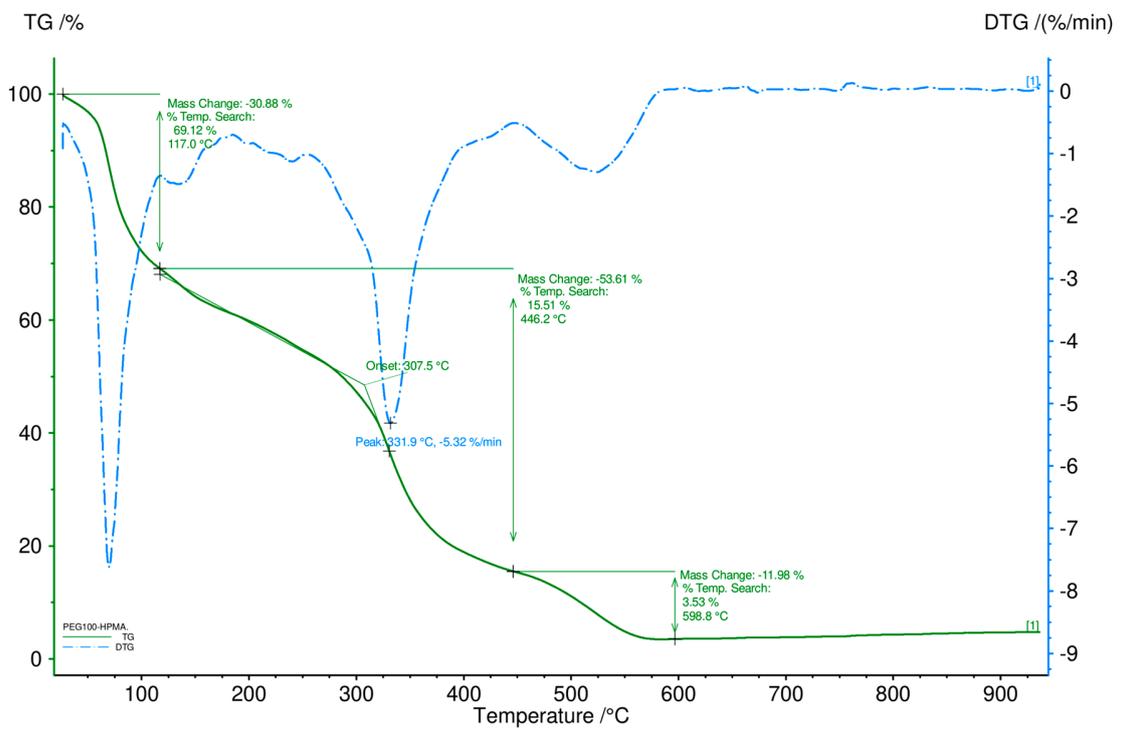


Figure S19. TG and DTG curves of PEG1000-HPMA.