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Supplementary Materials

Thermoresponsive Poly(*N*,*N*-diethylacrylamide-*co*-glycidyl methacrylate) Copolymers and Its Catalytically Active α-Chymotrypsin Bioconjugate with Enhanced Enzyme Stability

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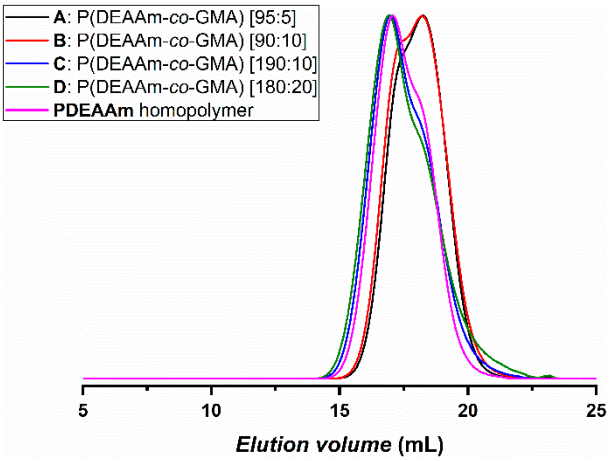

Figure S1. GPC chromatograms of the P(DEAAm-*co*-GMA) copolymers and PDEAAm homopolymer.

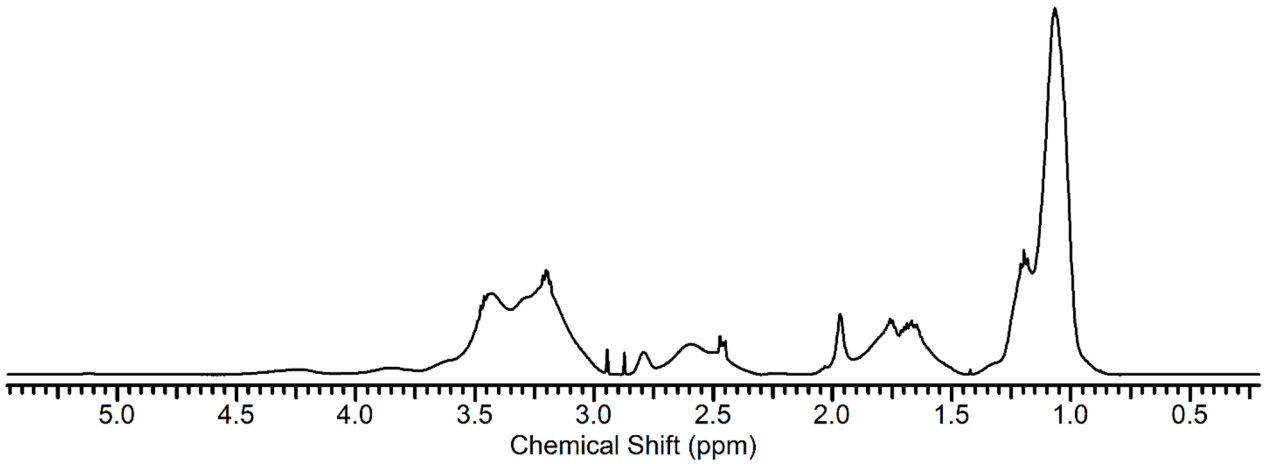

Figure S2. [1]H NMR spectrum of *Sample **B*** P(DEAAm-*co*-GMA) copolymer (molar feed ratio AIBN:DEAAm:GMA = 1:90:10).

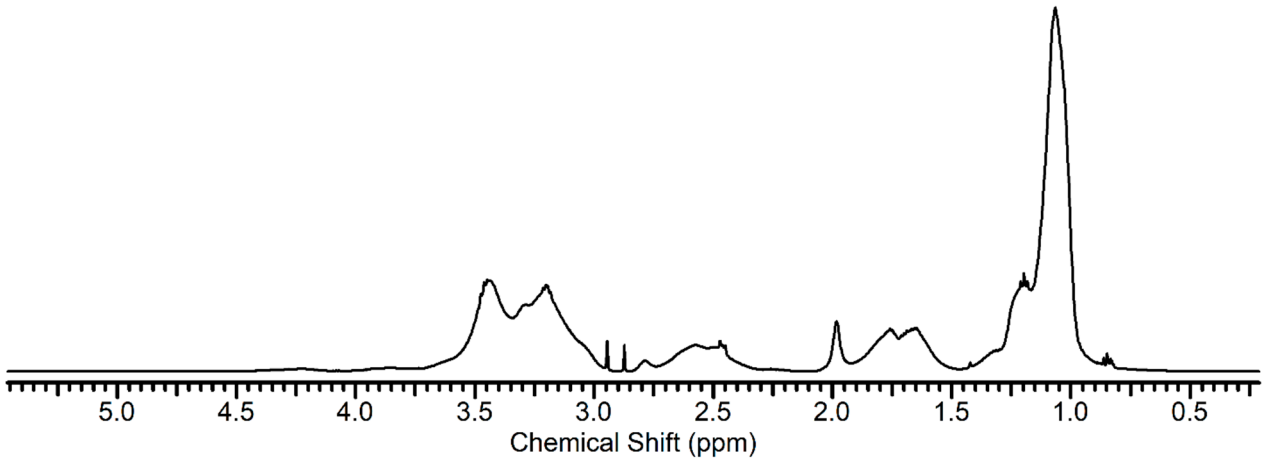

Figure S3. [1]H NMR spectrum of *Sample **C*** P(DEAAm-*co*-GMA) copolymer (molar feed ratio AIBN:DEAAm:GMA = 1:190:10).

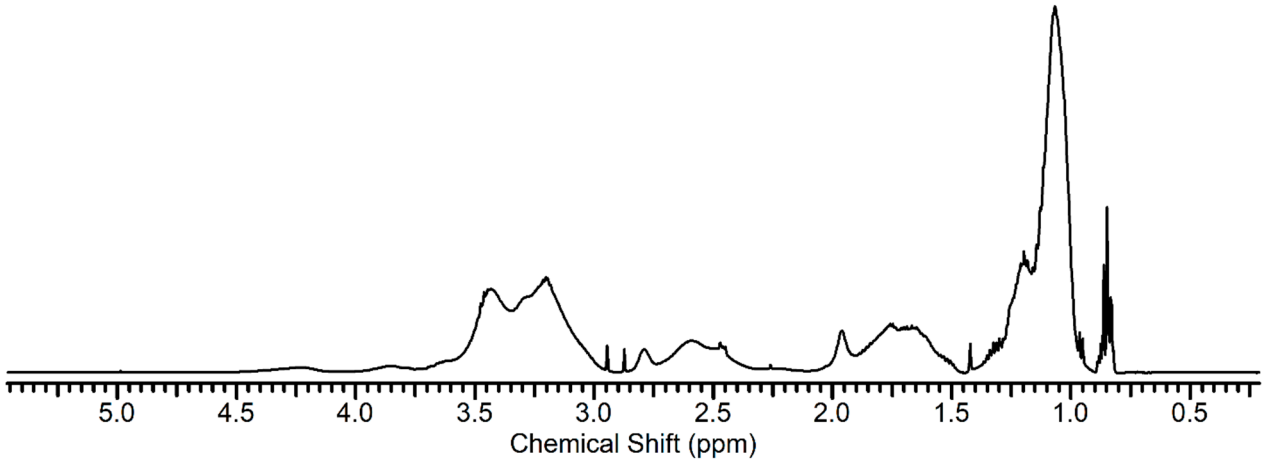

Figure S4. [1]H NMR spectrum of *Sample **D*** P(DEAAm-*co*-GMA) copolymer (molar feed ratio AIBN:DEAAm:GMA = 1:180:20).

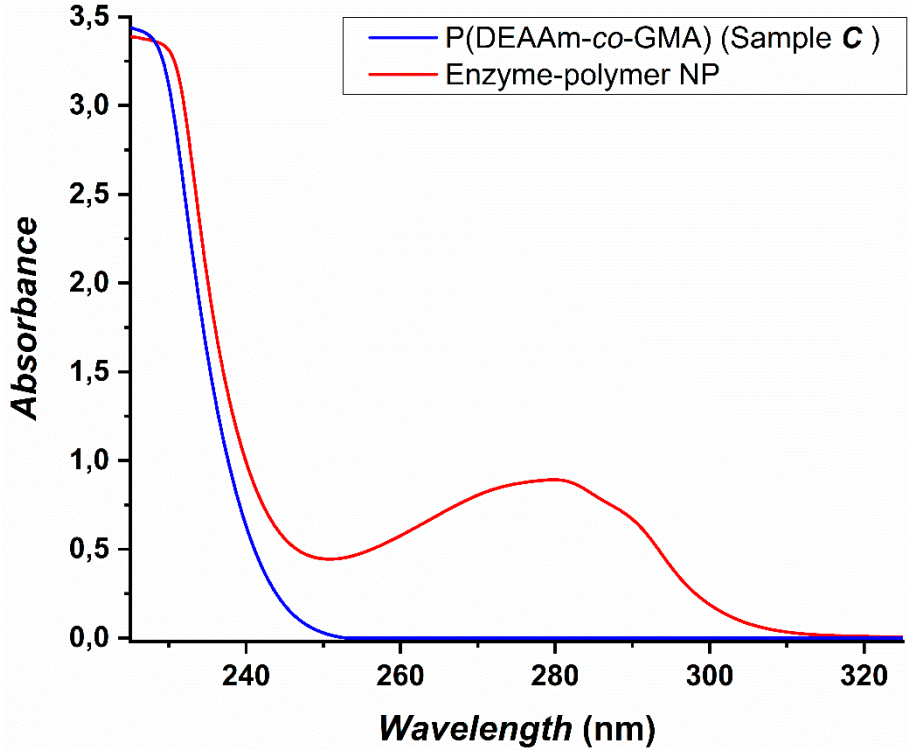

Figure S5. UV spectra of the P(DEAAm-*co*-GMA) (Sample ***C***, *blue*) and the produced enzyme-polymer nanoparticle (*red*).

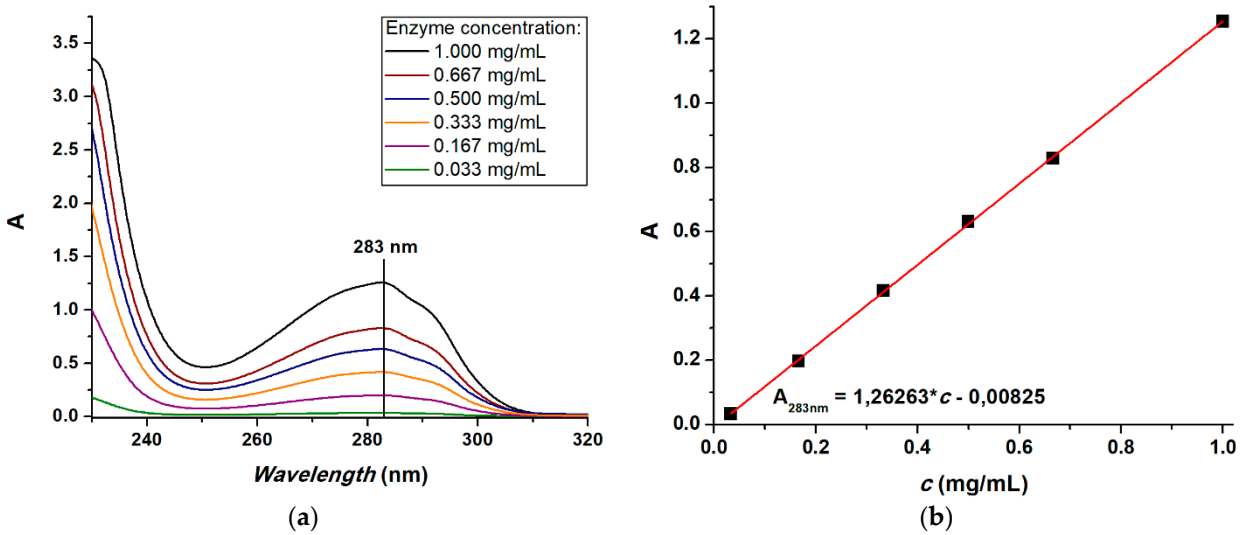

Figure S6. UV spectra of the α-chymotrypsin in the concentration range of 0.033-1 mg/mL (**a**) and the calibration curve fitted on the absorbance at 283 nm as a function of the enzyme concentration (**b**).

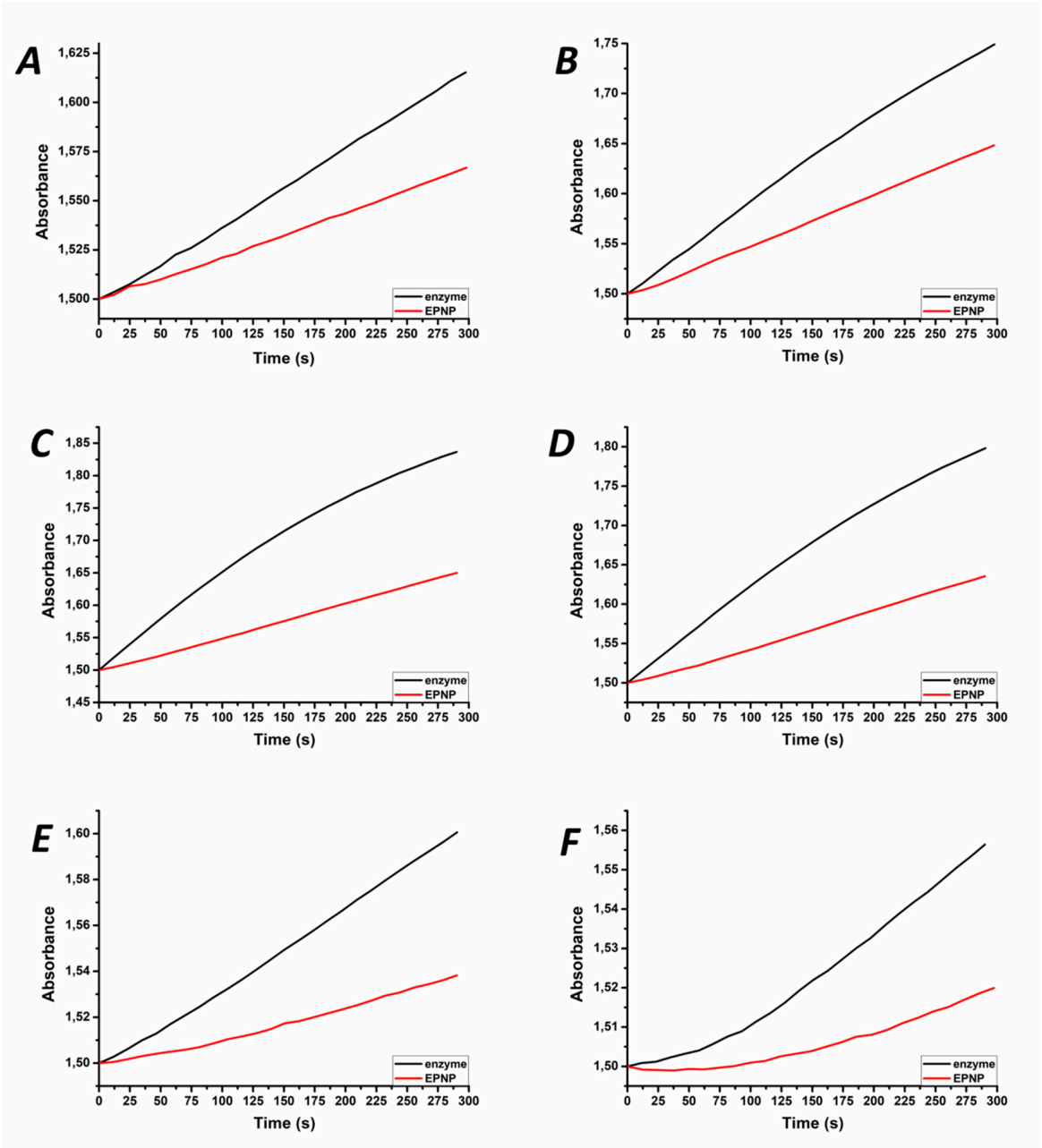

Figure S7. Representative enzymatic activity investigation curves of the absorbance measurement of the enzyme (black) and EPNP (red) in time in different pH solvents (pH = 6 (*A*); 7 (*B*); 7.4 (*C*); 7.8 (*D*); 8 (*E*); 9 (*F*)).

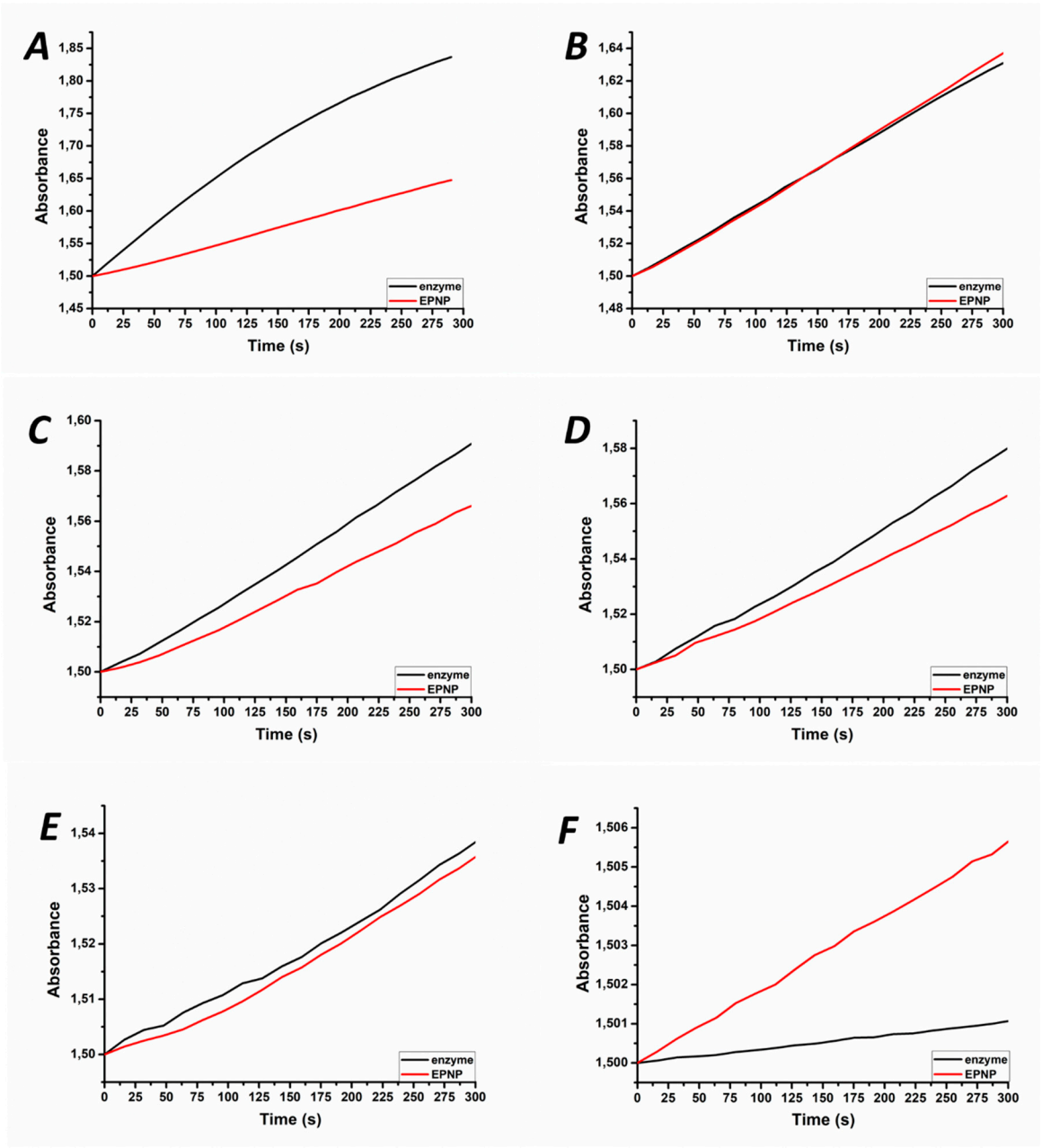

Figure S8. Representative curves of the activity measurements of the enzyme (black) and EPNP (red) in PBS buffer after thermostated at 45 °C for 0 min (***A***), 5 min (***B***), 15 min (***C***), 30 min (***D***), 60 min (***E***) and 120 min (***F***).