Supporting Information

Inspired by elastomers: Fabrication of hydrogels with tunable properties and re-shaping ability via photo-crosslinking at macromolecular level

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Fig. S1 ¹H NMR spectra of (a) DHBP and (b) DEBP





Fig. S2 Photographs of model compounds (left) and PEA5%-PAAm hydrogels (right) under different UV irradiation time



Fig. S3 Tensile stress–strain curves of (a) PEA-PAAm hydrogels with different content of PEA before UV irradiation, (b) TEMED hydrogels and TEMED-BIS hydrogels.

Sample	AAm/g	PEA/g	APS/g	TEMED/g	MBA/g	Water content	
PEA0.5%	10	0.05	0.01	0	0	70%	
PEA1%	10	0.1	0.01	0	0	70%	
PEA3%	10	0.3	0.01	0	0	70%	
PE5%	10	0.5	0.01	0	0	70%	
TEMED	10	0	0.01	0.053	0	70%	
TEMED-BIS	10	0	0.01	0.053	0.05	70%	

Table S1. Compositions of hydrogels including the conventional chemically cross-linked TEMED-BIS hydrogels using organic BIS as cross-linker and PEA-PAAm hydrogels with PEA as the cross-linker.

Table S2. The formulas of the model compounds.

Sample	AAm/g	PEA/g	APS/g	Water/g
PEA-APS-0.5%	0.5	0.5	0.0025	10
PEA-APS-1%	0.5	0.5	0.005	10

Table S3. Mechanical properties (Elongation at break, Tensile modulus and tensile strength) comparison of PEA-PAAm hydrogels with varied contents of PEA before and after UV irradiation 120 min.

Hydrogals	UV irradiation	Elongation at	Tensile modulus	Tensile strength			
Trydrogers	time (min)	break (mm/mm)	(kPa)	at break (kPa)			
PEA0.5%-PAAm	0	18.81	107	577			
	120	14.92	115	590			
PEA1%-PAAm	0	18.38	83	521			
	120	13.64	98	623			
PEA3%-PAAm	0	17.89	52	359			
	120	12.49	72	552			
PEA5%-PAAm	0	15.34	42	171			
	120	12.00	75	500			