Smart Low Molecular Weight Hydrogels with Dynamic Covalent Skeleton

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Experimental section

Synthesis of 3-(2-(4-formylphenoxy) ethyl)-1-methyl imidazolium bromide (MA).



Scheme S1. The synthesis route for MA.

The ¹H NMR (300 MHz, CDCl₃) of compound A: δ H / ppm 3.67 (t, 2H), 4.38 (t, 2H), 7.04 (d, 2H), 7.89 (d, 2H), 9.90 (s, 1H). ¹³C NMR (75.46 MHz, CDCl₃): δ C / ppm 31.53, 68.57, 115.51, 130.43, 132.30, 163.27, 191.80. MS: ([C₉H₉BrO₂+H]⁺) m/z = 228.9859, found m/z = 228.9836. The ¹H NMR (300 MHz, D₂O) of MA: δ H / ppm: 3.77 (s, 3H), 4.42 (t, 2H), 4.56 (t, 2H), 7.02 (d, 2H), 7.32 (s, 1H), 7.48 (s, 1H), 7.78 (d, 2H), 9.65 (s, 1H). ¹³C NMR (75.46 MHz, D₂O): δ C / ppm 35.80, 48.78, 66.27, 115.11, 122.73, 123.63, 129.73, 132.67, 163.22, 194.78. MS: ([C₁₁H₁₅N₂O₂]⁺) m/z = 232.21, found m/z = 231.1162.

Synthesis of 3, 3'-dithiobis (propionohydrazide) (DSPDZ).

¹H NMR (300 MHz, DMSO) of DSPDZ: δH / ppm: 9.07 (s, 1H), 4.22 (s, 2H), 2.89 (t, 2H), 2.42 (t, 2H).

Additional Results



Fig. S1 ESI-MS profile of the low molecular weight supra-gelator (BMBS).



Fig. S2 SEM image of BMBS hydrogel prepared from MA (100 mM) and

DSPDZ (50 mM).



Scheme S2. Chemical structures of CM and APDZ.



Fig. S3 Photographs of the solutions of MA and CM (a); APDZ (b).



Fig. S4 Optical microscope images of the self-healing process of BMBS hydrogels (MA, 100 mM and DSPDZ, 50 mM). 1) 0, b) 3, c) 6 and d) 12 hours, respectively. White arrows indicate the positions of cut gaps on the hydrogel samples. Scar bars: $100 \mu m$.



Fig. S5 Photographs of the resultant hydrogels by the precursor of MA and DSPDZ with different pH.



Fig. S6 Two disk-shaped hydrogels (the one on the left stained with

rhodamine B dye) were cut into 4 pieces and put together alternately. (b-e) Photos were taken after 0, 3, 6 and 12 hours.