Supporting Information

Thermo, pH and reduction responsive coaggregates comprising AB_2C_2 star terpolymers for multi-triggered release of doxorubicin

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Table S1 Dependence of apparent drug-release rate (K), cumulative release (CR) and increment of cumulative release (ICR) at 48 h on external stimuli during in vitro DOX release from various aggregates

Aggregates	stimuli	T (°C)	pН	DTT	<i>K</i> (h ⁻¹)	CR (%)	ICR(%)
S1	dual	25	7.4	none	0.0291	19.2	0
S 1	dual	25	7.4	yes	0.0705	41.4	116
S1	dual	37	7.4	none	0.115	37.0	93
S 1	dual	37	7.4	yes	0.161	61.1	218
S3	dual	37	7.4	none	0.065	26.7	0
S3	dual	37	7.4	yes	0.137	43.0	61
S 3	dual	37	5.3	none	0.119	38.4	44
S 3	dual	37	5.3	yes	0.215	63.1	136
S1+S3	triple	25	7.4	none	0.0261	18.3	0
S1+S3	triple	25	7.4	yes	0.0393	36.8	101
S1+S3	triple	25	5.3	none	0.0326	35.0	91
S1+S3	triple	25	5.3	yes	0.0496	42.0	130
S1+S3	triple	37	7.4	none	0.0618	24.1	32
S1+S3	triple	37	7.4	yes	0.135	42.9	134
S1+S3	triple	37	5.3	none	0.0837	33.0	80
S1+S3	triple	37	5.3	yes	0.177	57.1	212



Fig. S1 ¹H (top) and ¹³C (bottom) NMR spectra of DCP.



Fig. S2 ¹H (top) and ¹³C (bottom) NMR spectra of HCP.



Fig. S3 IR spectra of trifunctional agents DCP and HCP.



Fig. S4 IR spectra of alkyne-functionalized PCL (a), PCL-b-PNIPAM (b) and PCL-b-PtBA (c).



Fig S5. ¹H NMR spectrum of PEG(PCL)₂(PAA)₂ star copolymer in DMSO-*d*₆.



Fig S6. TEM images of copolymer aggregates formed by $PEG(PCL)_2(PNIPAM)_2$ (a, T = 25 °C; b and c, T = 37 °C), $PEG(PCL)_2(PAA)_2$ (d, T = 37 °C) and their mixtures according to equimolar ratio (e, T = 25 °C; f, T = 37 °C) in PBS solution with pH 7.4 (*c* = 0.50 mg mL⁻¹).



Fig. S7 DLS plots of S1/S3 coaggregates ($c = 0.50 \text{ mg mL}^{-1}$) in PBS solution (pH 7.4, 50 mM) at 25 or 37 °C for different time periods (t = 0 and 48 h).