

Supporting Information

Elaboration on the Distribution of Hydrophobic Segments in the Chains of Amphiphilic Cationic Polymers for siRNA Delivery

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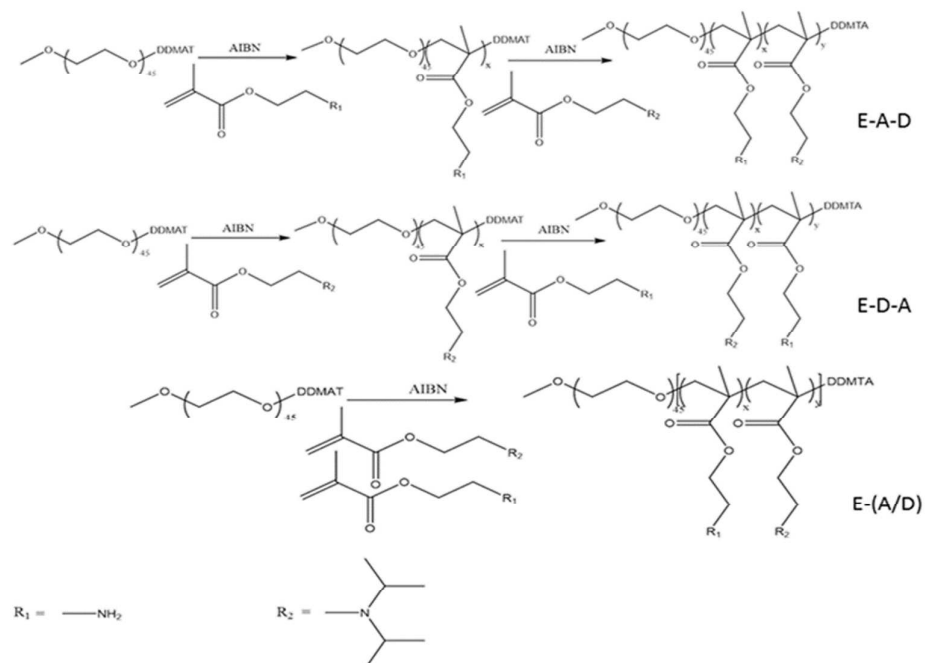
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Scheme S1. The synthetic routes of E-A-D, E-D-A and E-(A/D).

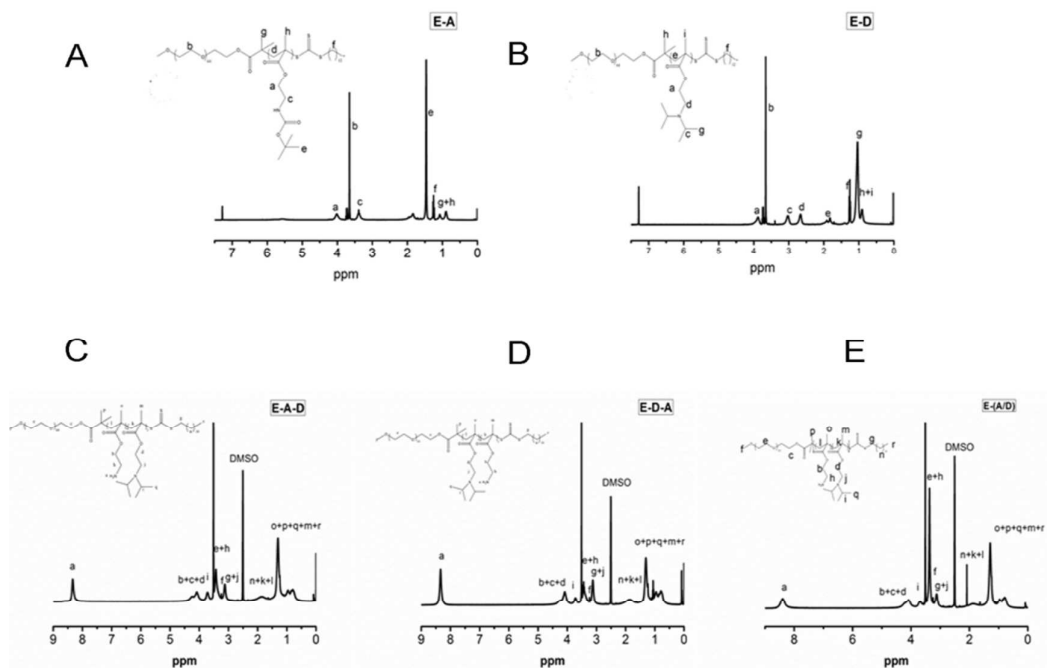


Figure S1. 1H -NMR spectroscopies of E-A, E-D, E-A-D, E-D-A and E-(A/D).

Table S1. The molecular structure, molecular weight and PDI of three polymers.

Sample	The molecular structure of the polymer ^a	M_n^b (10^4 g/mol)	PDI ^c
E-A-D	PEG ₄₅ -PAM ₄₆ -PDP ₄₅	2.21	1.67
E-D-A	PEG ₄₅ -PDP ₄₅ -PAM ₄₄	2.17	1.68
E-(A/D)	PEG ₄₅ -P(AM ₄₂ /DP ₄₂)	2.06	1.57

^a The molecular structure of the polymer was determined by ¹H NMR.

^b The mean molecular weight (M_n) of copolymers was estimated by ¹H NMR.

^c PDI= M_w/M_n measured by GPC.

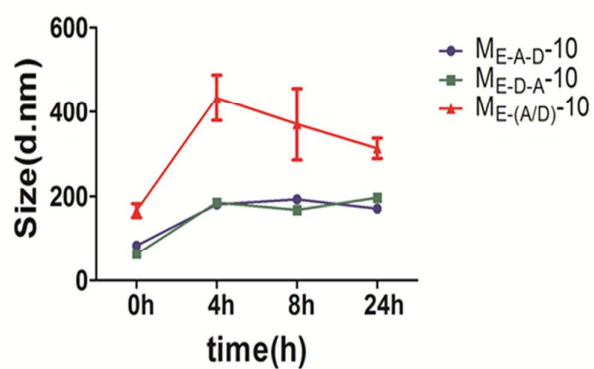


Figure S2. The stability of siRNA-loaded micelles at N/P=10 in PBS.

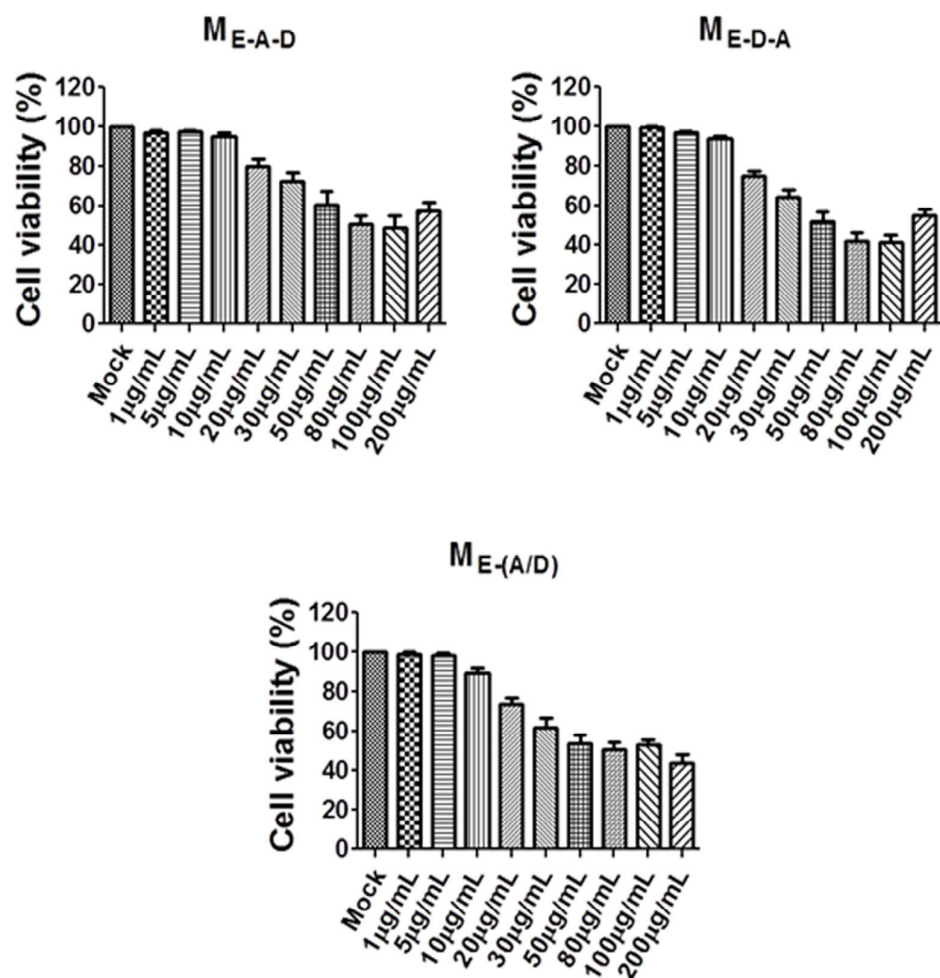


Figure S3. Cell cytotoxicity of M_{E-A-D}, M_{E-D-A} and M_{E-(A/D)} at different concentrations.

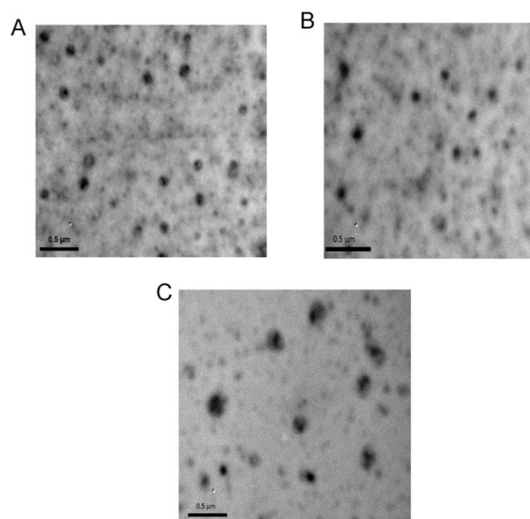


Figure S4. TEM images of M_{E-A-D} (A), M_{E-D-A} (B) and M_{E-(A/D)} (C).

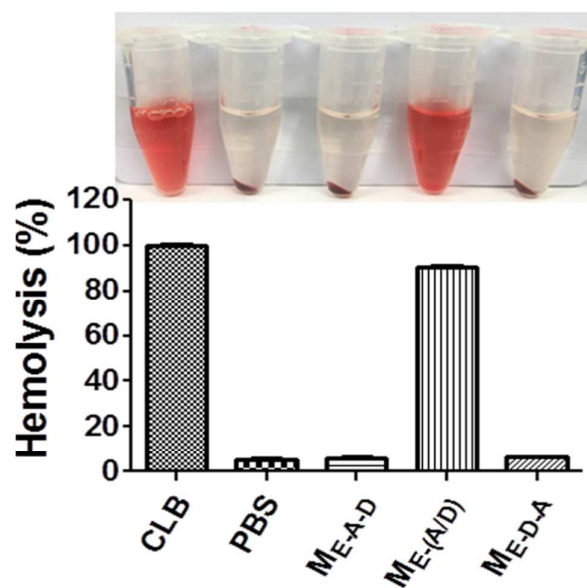


Figure S5. Hemolysis assay of 5 mM micelles in 1 mL PBS at pH=7.4. Cell lysis buffer was used as positive control and PBS was used as negative control.

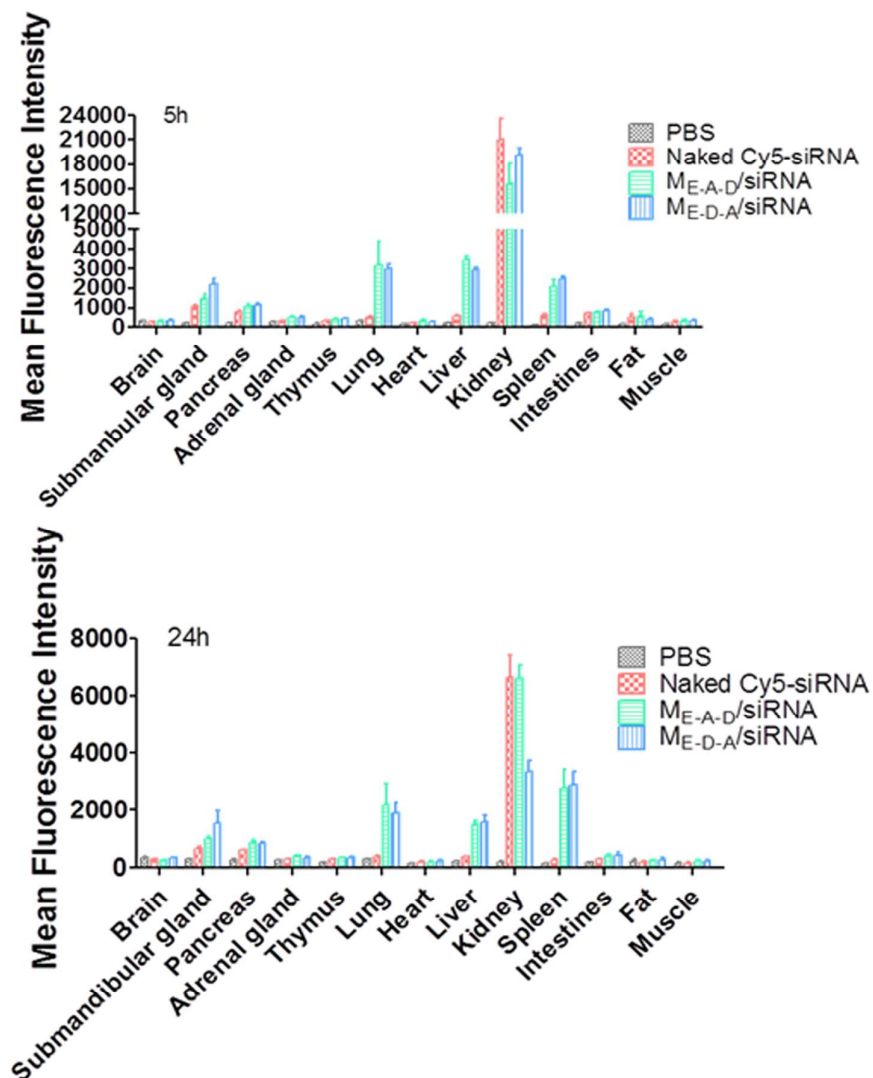


Figure S6. Quantitative analysis of different organs using a molecular imaging software package at 5 h and 24 h. Data were normalized to corresponding tissues from saline-treated animals. Each bar represents the mean±S.E. of three different experiments.

Table S2. The sequences of the siRNA used in this work.

siRNA Name	Sequences
Cy5-siRNA	sense: 5'-Cy5-CCUUGAGGCAUACUUCAAAdTdT-3'
	anti-sense: 5'-UUUGAAGUAUGCCUCAAGGdTdT-3'
siFL	sense: 5'-CCCUAUUCUCCUUCUUCGCdTdT-3'
	anti-sense: 5'-GCGAAGAAGGAGAAUAGGGdTdT-3'
siApoB	sense: 5'-GUCAUCACACUGAAUACCAAUdTdT-3'
	anti-sense: 5'-AUUGGUAAUUCAGUGUGAUGACACdTdT-3'
siPLK1	sense: 5'-UGAAGAAGAUACCCUCCUAdTdT-3'
	anti-sense: 5'-UAAGGAGGGUGAUCUUCUUCAdTdT-3'
siRRM2	sense: 5'-GCGAUUUAGCCAAGAAGUUCA-3'
	anti-sense: 5'-UAGCGACUAAACACAUCAAUU-3'