

Electronic Supplementary Information

Hydroxyl-containing non-viral lipidic gene vectors with macrocyclic polyamine headgroups

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Table S1. Particle size and zeta-potential of **6a** and **7a** against two weeks. The molar ratio of lipid/DOPE was 1 : 3.

Liposome	Particle size (nm)			Zeta-potential (mV)		
	Fresh liposome	7 days	14 days	Fresh liposome	7 days	14 days
6a	139.9 ± 3.4	139.7 ± 1.0	140.3 ± 2.6	52.5 ± 2.8	54.3 ± 1.7	49.0 ± 1.7
7a	134.6 ± 0.6	135.9 ± 0.5	134.4 ± 0.6	50.8 ± 0.5	52.6 ± 0.4	49.0 ± 0.7

Table S2. Particle size of liposome and lipoplexe against serum. “N/P = ∞” means that the liposomes without DNA. The molar ratio of lipid/DOPE was 1 : 3.

Liposome	N/P	With out serum (nm)	Serum (nm)
6a	∞	139.9 ± 3.4	259.3 ± 15.0
	6	163.8 ± 2.9	263.9 ± 9.8
6b	∞	139.9 ± 2.3	244.0 ± 1.9
	6	185.7 ± 19.9	258.5 ± 6.5
6b	∞	128.8 ± 2.7	227.8 ± 11.1
	6	165.5 ± 27.7	256.7 ± 23.5
6c	∞	120.5 ± 1.6	208.8 ± 19.6
	6	122.6 ± 9.7	208 ± 18.0
7a	∞	134.6 ± 2.2	257.4 ± 15.8
	6	155.5 ± 7.7	249.4 ± 11.0
7b	∞	142.8 ± 0.4	249.4 ± 8.2
	6	168.8 ± 5.3	233.2 ± 14.7

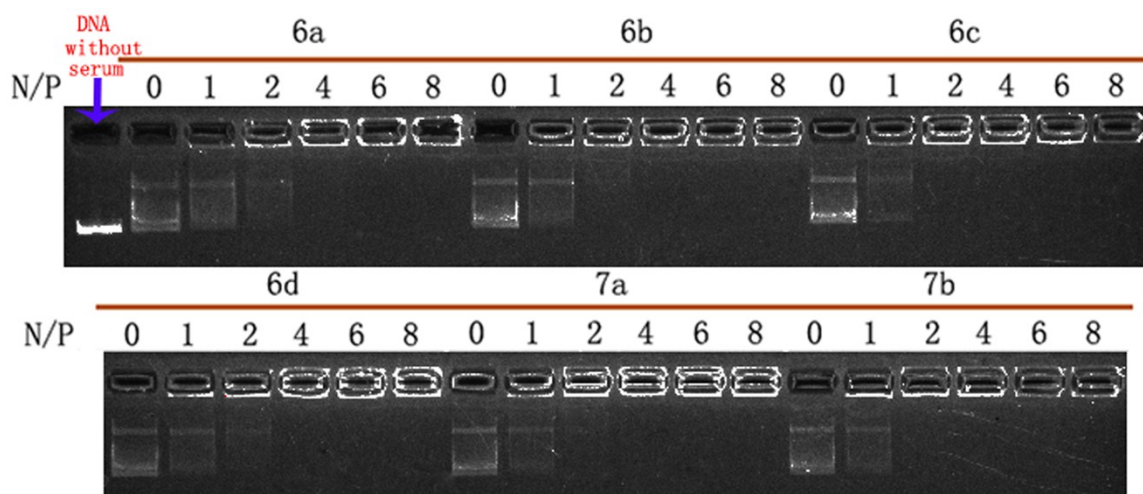


Fig. S1. Electrophoretic gel retardation assays of lipoplexes at different N/P ratios in the presence of 10% serum. The molar ratio of lipid/DOPE was 1 : 3.

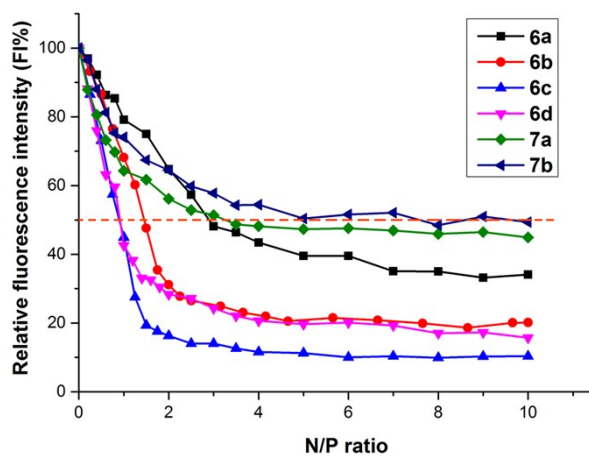


Fig. S2. Fluorescence quenching of EB by lipids 6 and 7/DOPE at various N/P ratios in 10 mM of HEPES buffer. The molar ratio of lipid/DOPE was 1 : 3.

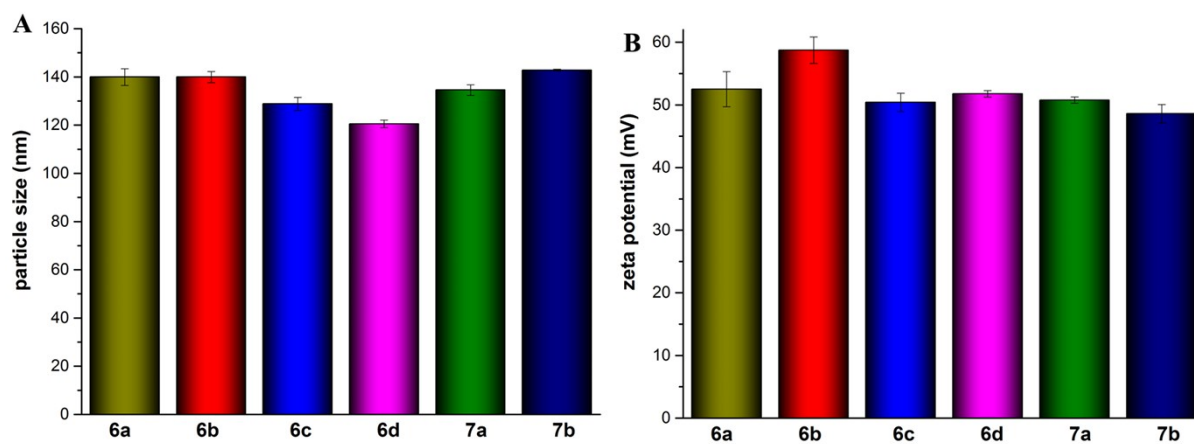


Fig. S3. Mean particle size (A) and Zeta-potential (B) of the liposomes without DNA (DLS at room temperature). The molar ratio of lipid/DOPE was 1 : 3.

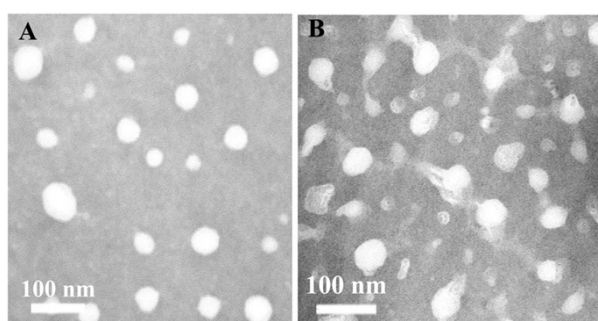


Fig. S4. TEM images of the liposomes formed from 6a (A) and 7a (B) in deionized water. The molar ratio of lipid/DOPE was 1 : 3.

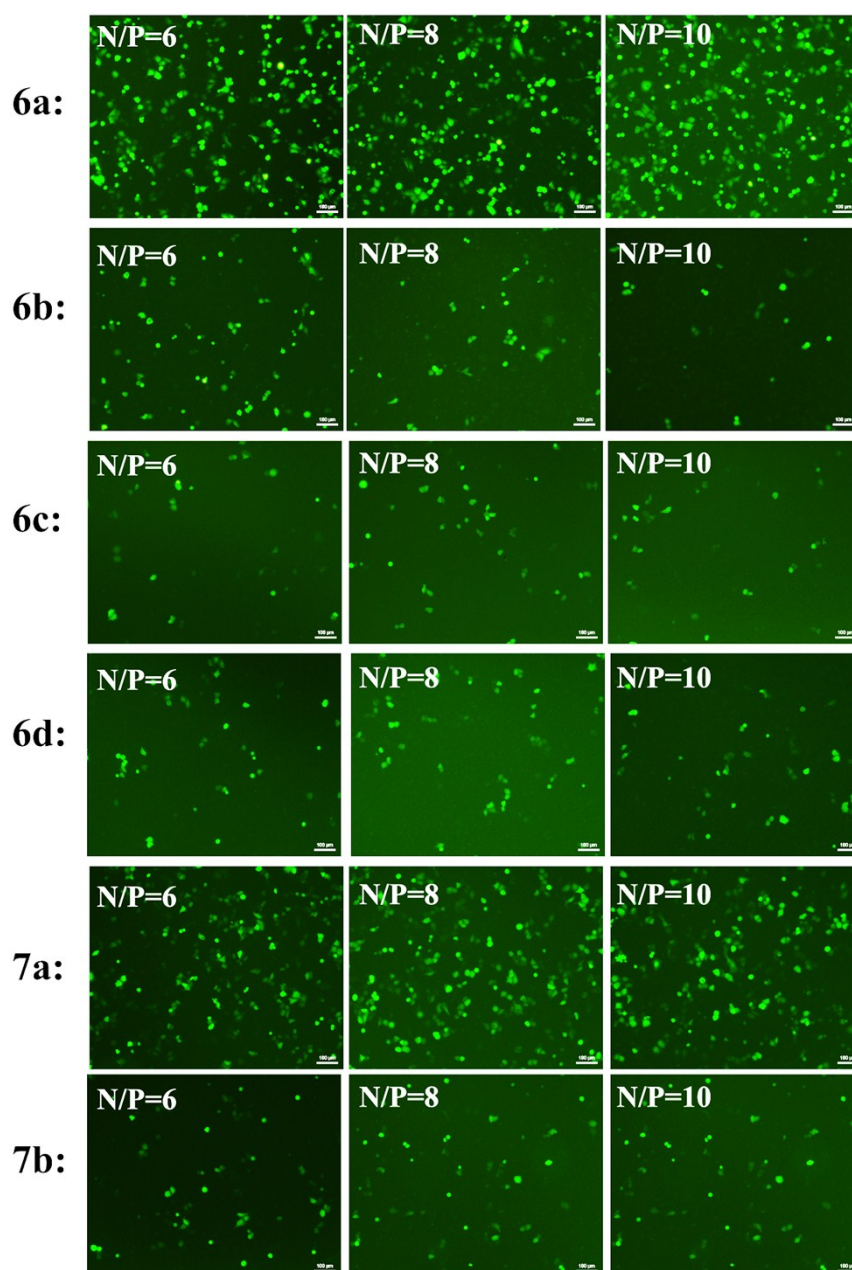


Fig. S5. Fluorescent microscope images of 7402 cells transfected by the lipoplexes formed from the six lipids. The cells were observed by fluorescence microscopy 24 h after transfection. The molar ratio of lipid/DOPE was 1 : 3.

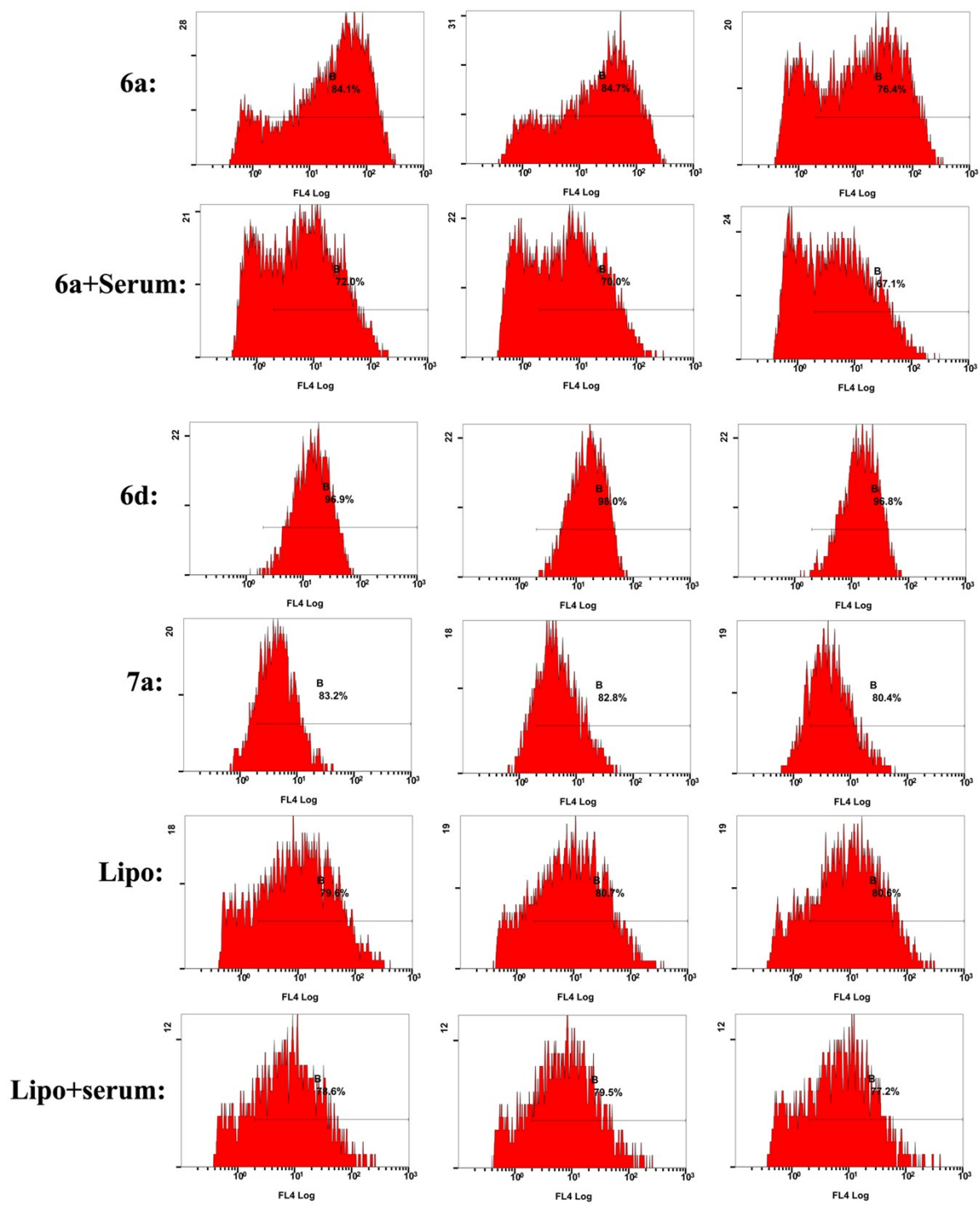


Fig. S6. Flow cytometry analysis of the liposomes at the optimal N/P ratio (N/P = 6 for **6a**, N/P = 8 for **6a** with serum, N/P = 8 for **6d** and N/P = 10 for **7a**). The molar ratio of lipid/DOPE was 1 : 3.