

***In vivo* siRNA delivery with dendritic poly(L-lysine) for the treatment of hypercholesterolemia**

Supplemental Information

Kazuto Watanabe,^a Mariko Harada-Shiba,^b Akira Suzuki,^b Risa Gokuden,^a Ryohsuke Kurihara,^a Yusuke Sugao,^a Takeshi Mori,^{ac} Yoshiki Katayama^{ac} and Takuro Niidome^{*acd}

^a Department of Applied Chemistry, Faculty of Engineering, Kyushu University, Fukuoka 819-0395, Japan. Fax & Tel: +81-092-802-2851; E-mail: kazuto@mail.cstm.kyushu-u.ac.jp; niidome.takuro.655@m.kyushu-u.ac.jp

^b Department of Bioscience, National Cardiovascular Center Research Institute, Suita, Osaka 565-8565, Japan. E-mail: mshiba@ri.ncvc.go.jp

^c Center for Future Chemistry, Kyushu University, Fukuoka 819-0395, Japan.

^d PRESTO, Japan Science and Technology Corporation, Kawaguchi, 332-0012, Japan.

† This article is part of a Molecular BioSystems ‘Emerging Investigators’ issue highlighting the work of outstanding young scientists at the chemical- and systems-biology interfaces.

Table S1 Effects of C/A ratio on the complex size and its ζ -potential. Ten $\mu\text{g/ml}$ double-stranded DNA (dsDNA) and KG6 at various C/A ratios were mixed in 700 μl of sterile solution containing 5% dextrose and 0.5 mM sodium chloride. The sizes and ζ -potentials of each complex were measured with a Zetasizer Nano ZS (Malvern Instruments Ltd., Worcestershire, United Kingdom). The dsDNA had the following sequences: 5'-GTCATCACA CTGAATACCAAT-3', 5'-ATTGGTATT CAGTGTGATGACAC-3', which was the same to si-ApoB^I sequence with emplacements of Us with Ts).

	C/A ratio of dsDNA/KG6 complex						
	0	0.5	1.0	2.0	4.0	8.0	16.0
Size (nm)	n.d.	151.7 $\pm$ 0.4	176.5 $\pm$ 7.9	185.2 $\pm$ 6.7	145.8 $\pm$ 2.8	168.4 $\pm$ 9.9	159.8 $\pm$ 6.1
ζ -potential (mV)	n.d.	-23.0 $\pm$ 0.3	16.6 $\pm$ 0.4	29.6 $\pm$ 0.9	27.8 $\pm$ 1.0	30.4 $\pm$ 0.9	25.2 $\pm$ 0.9

Data are means $\pm$ S.E.

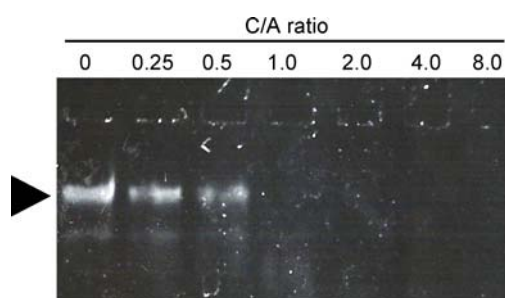


Fig. S1 Effects of C/A ratio on binding ability of KG6 with siRNA. si-ApoB^I (200 ng) and KG6 at various C/A ratios were mixed in 10 μl of sterile water, and then the complexes were applied to 20% (w/v) native polyacrylamide gel in TBE (45 mM Tris-Borate and 1 mM EDTA, pH 8.0) buffer. After 60 min electrophoresis at 100 V, the gel was stained with SYBR Gold. The arrowhead on the left of the gel indicated position of the naked siRNA.