

Supporting Information

Plasmid DNA delivery using fluorescein-labeled arginine-rich peptides

Makoto Oba^{a,*}, Yosuke Demizu^b, Hiroko Yamashita^{b,c}, Masaaki Kurihara^{b,c}, Masakazu Tanaka^a

^aGraduate School of Biomedical Sciences, Nagasaki University, Nagasaki 852-8521, Japan

^bDivision of Organic Chemistry, National Institute of Health Sciences, Tokyo 158-8501, Japan

^cGraduate School of Bioscience and Biotechnology, Tokyo Institute of Technology, Yokohama 226-8501, Japan

*To whom correspondence should be addressed:

Makoto Oba

Graduate School of Biomedical Sciences, Nagasaki University, Nagasaki 852-8521, Japan

Tel: +81-95-819-2424; Fax: +81-95-819-2424

E-mail: moba@nagasaki-u.ac.jp (M. Oba)

Peptide purity was checked by analytical HPLC using Discovery® BIO Wide Pore C18 column (25 cm x 4.6 mm).

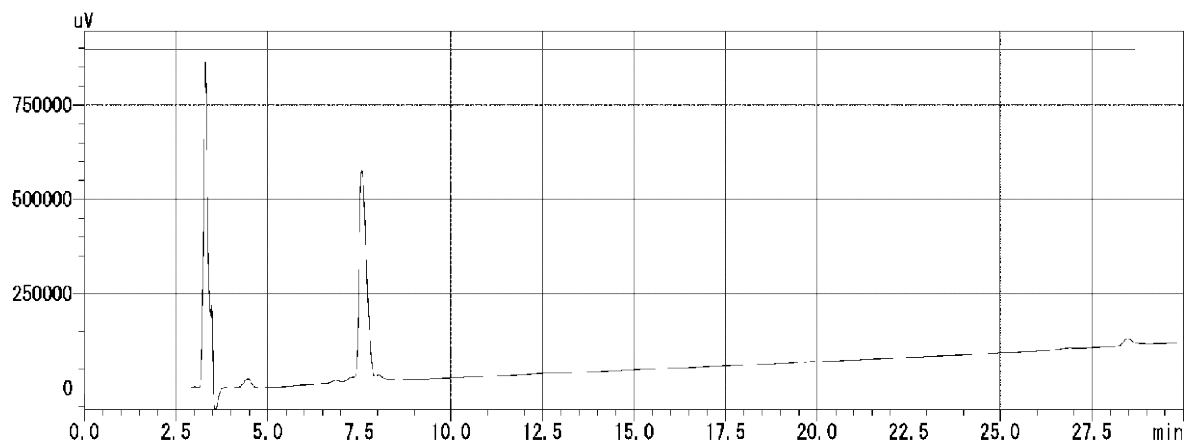
solvent A; 0.1% TFA/water

solvent B; 0.1% TFA/MeCN

flow rate; 1.0 mL•mL⁻¹

gradient; 10-60% gradient of solvent B over 30 min

6-FAM-β-Ala-(L-Arg-L-Arg-Gly)₃-NH₂ [(RRG)₃]



ESI(+)-TOF :

m/z calcd for C₄₄H₈₆N₂₈O₁₀ [M+H]⁺ 1167.72

m/z	calc.	Found
[M+2H] ^{+2/2}	584.87	584.62
[M+3H] ^{+3/3}	390.25	389.85
[M+4H] ^{+4/4}	292.94	292.77