Supplementary Information

Preparation of siRNA-PLGA/Fab'-PLGA mixed micellar system with target cell-specific recognition

Mai Hazekawa^{1*}, Takuya Nishinakagawa¹, Takeshi Mori², Miyako Yoshida², Takahiro Uchida² and Daisuke Ishibashi¹

¹Department of Immunological and Molecular Pharmacology, Faculty of Pharmaceutical Sciences, Fukuoka University, Fukuoka, Japan

²Department of Clinical Pharmaceutics, Faculty of Pharmaceutical Sciences, Mukogawa Women's University, Hyogo, Japan



F(ab')2 and Fab' structure obtained by fragmentation of IgG and existence confirmation and their activity evaluation for the antigen

(a) Shape of the fragmentation of IgG; (b) Detection of IgG fragmentation using SDS-PAGE; (c) Evaluation of the binding activity of antibody fragments used in this study using a flow cytometer.



Characterization of siRNA-PLGA and Fab'-PLGA conjugate via covalent bond (a) Fab' visualized by Ag staining in the gel after SDS-PAGE. Lane1: F(ab')2 and PLGA(FITC) mixture, Lane2: Fab'-PLGA(FITC), Lane3: Fab'-PLGA(FITC) treated with DTT (b) FITC visualized by transilluminator in the gel after SDS-PAGE. Lane1: F(ab')2 and PLGA(FITC) mixture, Lane2: Fab'-PLGA(FITC), Lane3: Fab'-PLGA(FITC) treated with DTT.



The size distribution of each conjugate micelles. (a) siRNA–PLGA conjugate micelles (b) siRNA–PLGA/Fab'–PLGA mixed micelles (c) Fab'–PLGA conjugate micelles.



S1b

S2a, Ag stein of SDS-PAGE

S2b, Fluorescence visualized by transilluminator of SDS-PAGE



Supplementary Figure S4 (continued)





Fig. 2b



Fig. 2c



Supplementary Figure S4 (continued)

Fig. 4a, CyclinB1



Supplementary Figure S4 (continued)

Fig. 4b, GAPDH

Exposure time 10 sec

