Supporting information

PEGylated Cationic Hybrid Bicellar Nanodisc for Efficient siRNA

Delivery

Yanyan Li^{a+}, Yidi Wu^{c+}, Shuquan Zheng^c, Xiaolong Liang^e, Xiaorui Han^c, Renfa Liu^d, Deyao Zhao^c, Yunhui Zhao^f, Yushen Jin^d, Min Chen^d, Xiaoxia Wang^c, Huiqing Cao^c, Xiuli Yue^{b+}, Tiejun Sten Shi^{a, g*}, Zicai Liang^{c*+}

^a School of Life Science and Technology, Harbin Institute of Technology, Harbin 150001, PR China, ^b School of Municipal and Environmental Engineering, Harbin Institute of Technology, Harbin 150001, PR China,

^{c.}Laboratory of Nucleic Acid Technology, Institute of Molecular Medicine, Peking University, Beijing 100871, PR China,

^{d.}Department of Biomedical Engineering, College of Engineering, Peking University, Beijing 100871, PR China,

^{e.}Department of Ultrasonography, Peking University Third Hospital, Beijing 100191, PR China

^{f.}School of Chemistry and Chemical Engineering, Hunan University of Science and Technology, Xiangtan 411201, PR China.

^{g.}Department of Biomedicine, University of Bergen, Bergen, Norway.

⁺These authors contributed equally to this work.

X.L.Yue: xiulidx@163.com

T. Shi: tiejun.shi@uib.no

Z.C.Liang: liangz@pku.edu.cn

Figure S1. Cytotoxicity of NDs. Viability of HepG2-Luc cells after 24h exposure to ND1 and ND5 by MTT assay.



Figure S2. Intracellular distributions and celluar uptake of ND1-4/siRNA complexes. (A) Intracellular distributions of Cy5-labeled siRNA loaded by the NDs were examed by confocal laser scanning microscopy (CLSM). LysoTracker®Greenwas used to stain lysosome (green), Hoechst 33342 was used to stain nuclei (blue) and siRNA was labelled with Cy5 (red).

