Electronic Supporting Information

Polyphenol-modified Nanovesicles for Synergistically Enhanced *in vitro* Tumor Cell Targeting and Apoptosis

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Supplementary data



Figure S1. (a) Size and zeta potential of NVs varying with stoichiometric ratio of DPPC/DC-Cholesterol. (b) Hydrodynamic diameters of the NVs. (c) TEM image of NVs.



Figure S2. (a) Synthesis of TAMA. (b) ¹H NMR of N-3-bromopropylmaleimide. (c) ¹³C NMR of TA and TAMA.



Figure S3. (a) Size and zeta potential of NV_{BC71} varying with the concentration of BC71. (b) Hydrodynamic diameter of $NV_{BC71-30 \ \mu M}$.



Figure S4. Fluorescence spectra of Texas red DHPE-loaded TANV_{BC71} after incubation with FBS/PBS (1/9, v/v) as a function of time.



Figure S5. (a) Fluorescence images of HaCaT cells treated with NV_{BC71-30 μ M, TA₅₀NV_{BC71-30 μ M} and TA₁₀₀NV_{BC71-30 μ M.} The scale bar is 50 μ m. (b) Cell viability (HaCaT) after treatment of NV_{BC71-30 μ M, TA₅₀NV_{BC71-30 μ M} or TA₁₀₀NV_{BC71-30 μ M} through live/dead fluorescence image analysis.}}



Figure S6. Quantitative cellular uptake of (a) TANVs with varying the concentration of conjugated TAMA (*p < 0.05 compared to treated NVs), (b) NV_{BC71} with varying the concentration of BC71 (*p < 0.05 compared to treated NVs), (c) TANV_{BC71-30 μ M} with varying the concentration of TAMA (*p < 0.05 compared to treated NV_{BC71-30 μ M}), and (d) TA₅₀NV_{BC71} with varying the concentration of BC71 (*p < 0.05 compared to treated TA₅₀NVs). HaCaT cells (black) and HCT116 cells (red) were used.