

Supporting Information

Immunogenic dead cell engineered by sequential treatments of ultraviolet irradiation/cryo-shocking for lung-targeting delivery and tumor vaccination

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Figure S1. CRT expression of Lewis cells after UV irradiation. Data were shown as mean \pm S.D. ($n = 6$).

Figure S2. Viability of Lewis cells after UV irradiation of different exposure doses. Data were shown as mean \pm S.D. ($n = 5$).

Figure S3. Cell morphology after UV irradiation at indicated irradiation intensities. Scale bar, 20 μm .

Figure S4. Particle size of HCPT/Lip and QS-21/Lip in PBS after storing at 4°C Data were shown as mean \pm S.D. ($n = 3$).

Figure S5. SEM image of HCPT&QS-21/UV-Cryo cell. Scale bar, 2 μm .

Figure S6. Ki67 and TUNEL staining assay of lungs. Scale bar, 100 μm .

Table S1. Hydrodynamic size, ζ -potential, entrapment efficiency (EE) and drug loading (DL) of HCPT-loaded liposomes. Data were shown as mean \pm S.D. ($n = 3$).

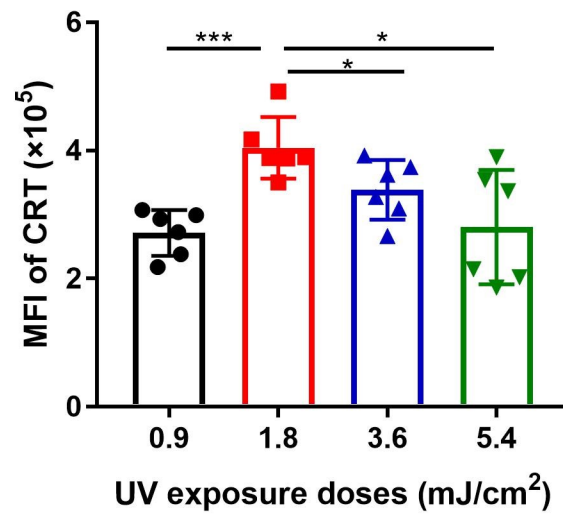


Fig. S1 CRT expression of Lewis cells after UV irradiation. Data were shown as mean \pm S.D. ($n = 6$). * $P < 0.05$ and *** $P < 0.001$.

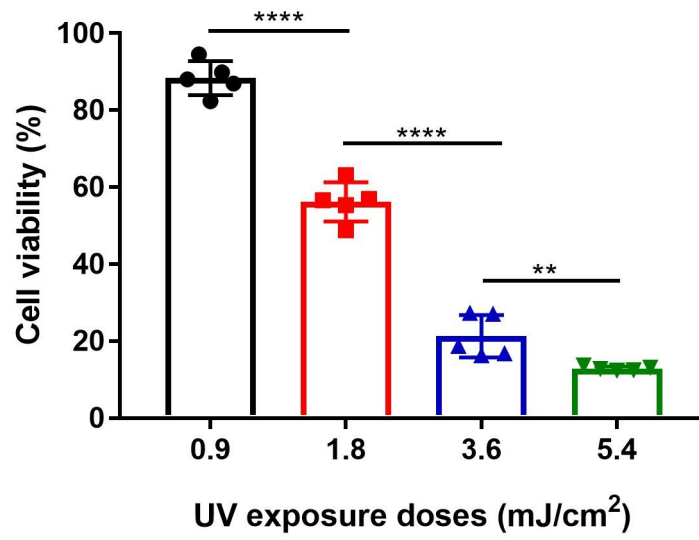


Fig. S2 Viability of Lewis cells after UV irradiation of different times. Data were shown as mean \pm S.D. ($n = 5$). ** $P < 0.01$ and **** $P < 0.0001$.

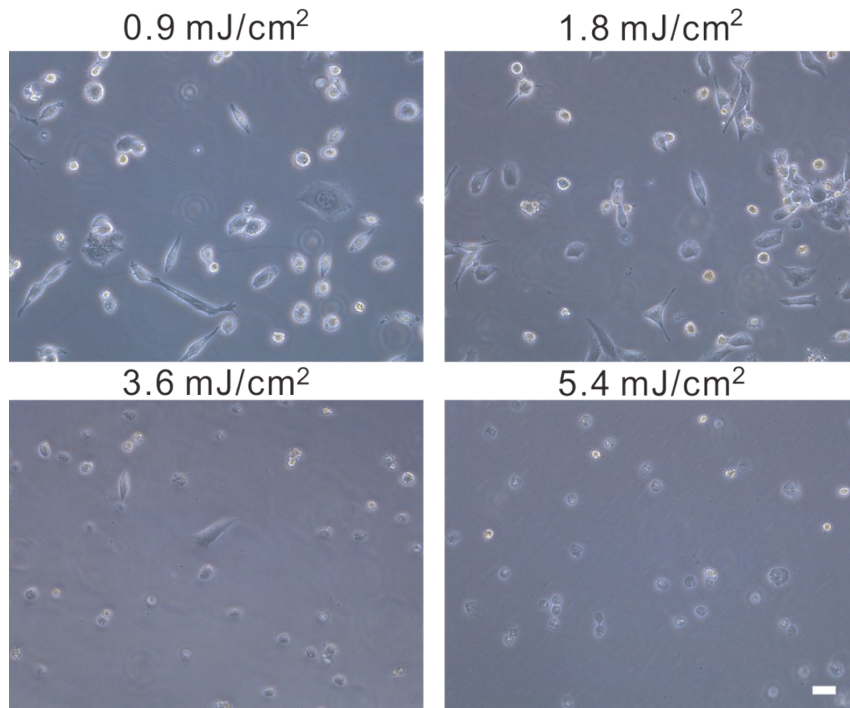


Fig. S3 Changes of cell morphology after UV irradiation at indicated irradiation intensities. Scale bar, 20 μm .

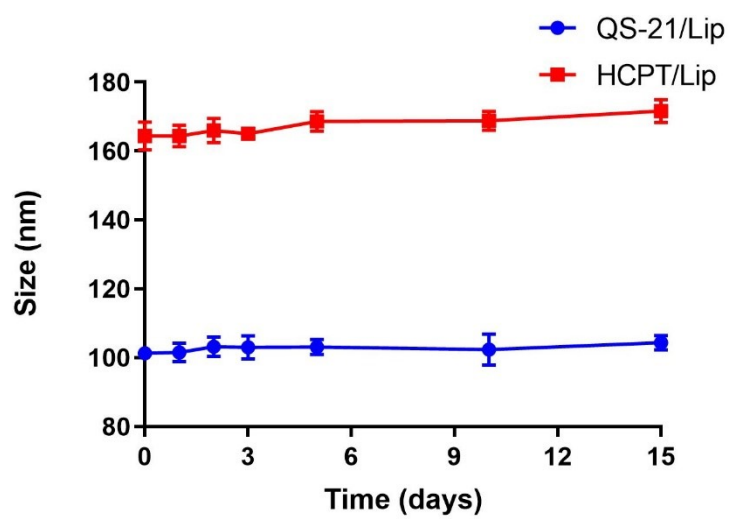


Fig. S4 Particle size of HCPT/Lip and QS-21/Lip in PBS after storing at 4°C. Data were shown as mean \pm S.D. ($n = 3$).

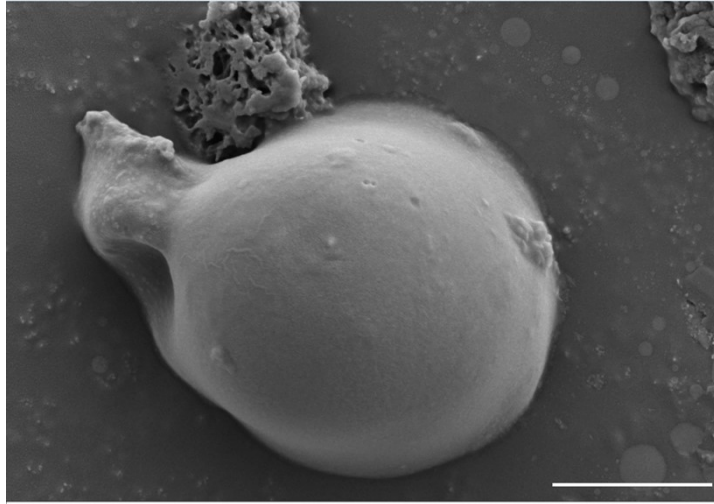


Fig. S5 SEM image of HCPT&QS-21/UV-Cryo cell. Scale bar, 2 μm .

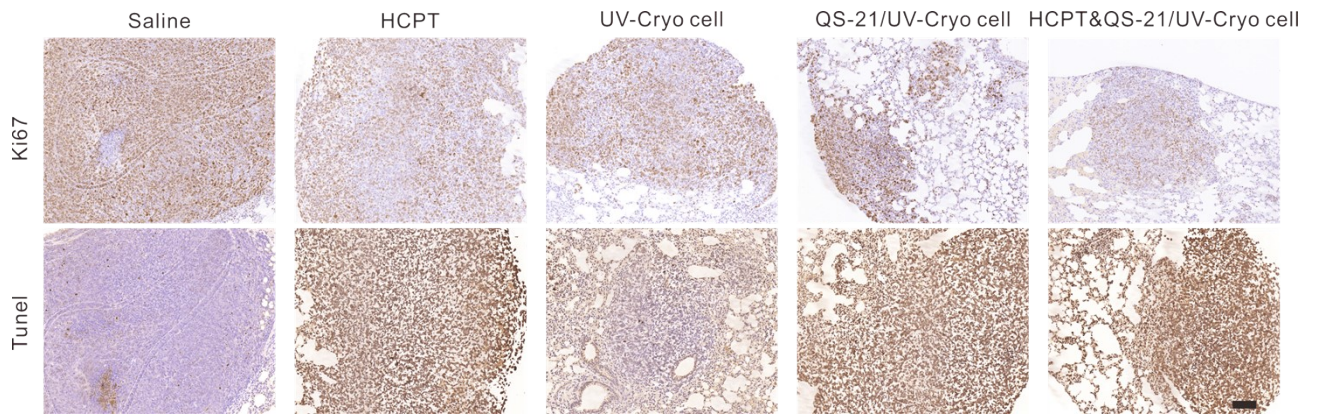


Fig. S6 Ki67 and TUNEL staining assay of lungs. Scale bar, 100 μ m.

Tab. S1 Physical characterizations of HCPT/Lip ($n = 3$).

Size (nm)	ζ -potential (mV)	EE (%)	DL (%)
163.7 ± 4.1	19.9 ± 0.7	68.2 ± 3.5	6.2 ± 0.3