

Supplemental information

Bioorthogonal oncolytic-virus nanovesicles combined bio-immunotherapy with CAR-T cell for solid tumors

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Supplemental figures:

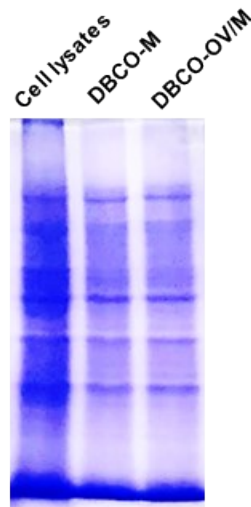


Figure S1. Sodium dodecyl sulfate - polyacrylamide gel electrophoresis (SDS-PAGE) analysis of cell lysates, DBCO-M, and DBCO-OV/M.

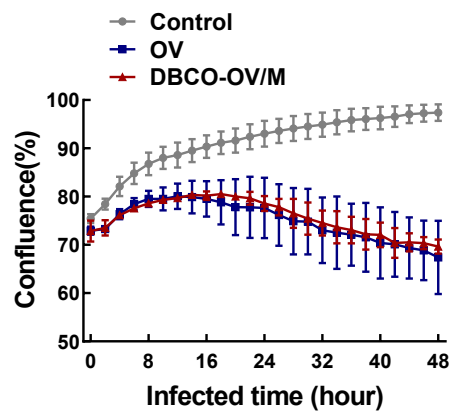


Figure S2. Cell confluence of SPCA-1 during the infection of OV or DBCO-OV/M in 48 h. The cytotoxicity of naked OV (1×10^8 VPs) and OV/M was analyzed by cell confluence. Cell confluence indicated host cell viability which was real-time monitored using Agilent xCELLigence RTCA.

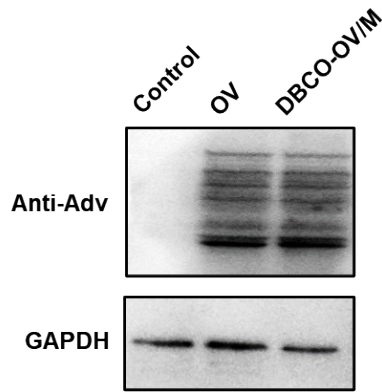


Figure S3. Western blotting assay of adenovirus proteins on OV or DBCO-OV/M treated cells, confirming the infecting capability of DBCO-OV/M.

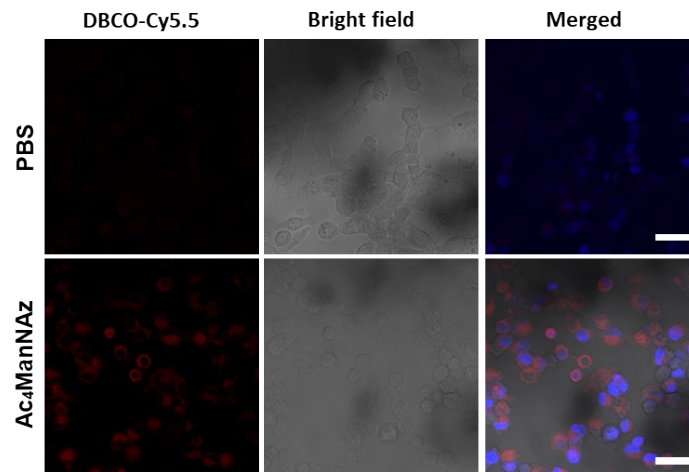


Figure S4. CLSM images of SPCA-1 cells treated with PBS or AC₄ManNAz. Azide modification for SPCA-1 cells through bioorthogonal glycometabolic labeling. DBCO-cy5.5 probe was used for detecting labeling of N₃ group. Scale bar= 25 μm.

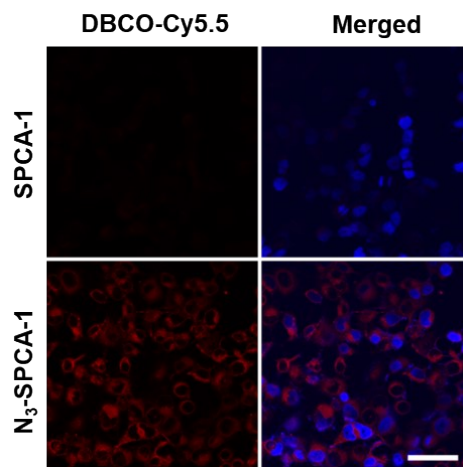


Figure S5. SPCA-1 subcutaneous tumor was injected with AC₄ManNAz for Azide modification. CLSM images of tumor slides of SPCA-1 or N₃-SPCA-1 treated with DBCO-Cy5.5. Scale bar= 100 μm.

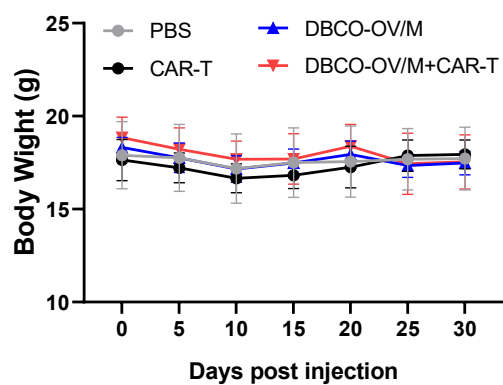


Figure S6. The body weight of mice treated with PBS (control), DBCO-OV/M nanovesicles, CAR-T, or DBCO-OV/M nanovesicles + CAR-T.

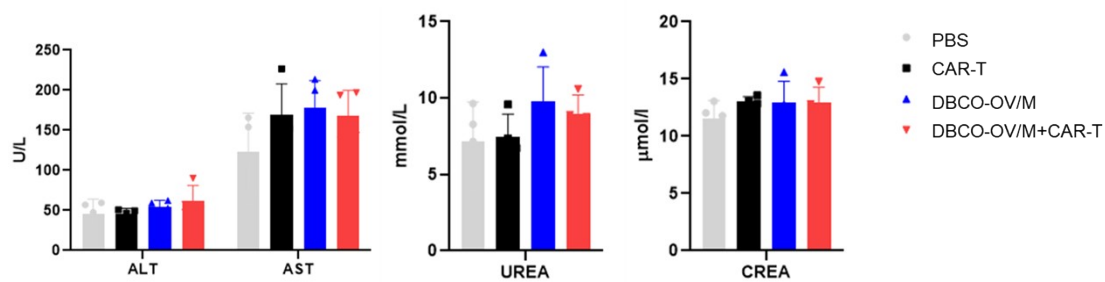


Figure S7. Liver and kidney functionality by blood test parameters after administration (mean \pm s.e.m., n = 3). no significance.

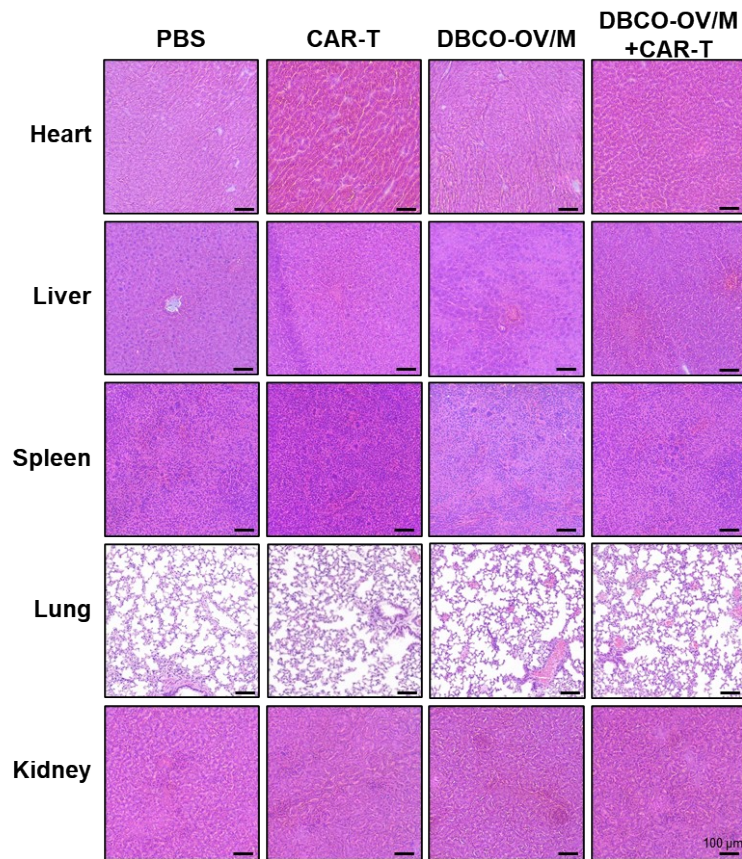


Figure S8. Tissue slices of major organ analyzed by Hematein-Eosin stain. No obvious differences in various groups were observed when compared with that in the PBS group. Scale bar = 100 μ m.