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SUPPORTING INFORMATION

Metabolic labelling of cancer cells with glycodendrimers stimulate immune-mediated cytotoxicity.

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Figure S 1. Flow cytometry analysis of BT-549 metabolically labelled with various concentrations of Ac₄ManNAz, followed by SPAAC conjugation to 50 μ M DBCO-PEG₄-Fluor 545. As control, cells without any treatment and non azido-labelled cells were used.



Figure S 2. DBCO-ARM1, 4 and 16 coupling to azido groups exposed at the BT-549 cell-surface. Azido labelled cells or unlabelled cells were treated with DBCO-ARMs or DBCO-PEG (10-0,1 μ M). SPAAC coupling of the DBCO-conjugates was revealed using anti-Rha IgM naturally present in HS and fluorescent secondary antibody. Cell fluorescence was analysed by flow cytometry. Data are presented as mean \pm SD (n = 3).



Figure S 3. Dose-response curve of the fluorescence of BT-549 cells metabolically labelled with the azido sugar Ac₄ManNAz, followed by SPAAC conjugation to DBCO-ABM16 at various concentrations (0 μ M, 0,1 μ M, 0,5 μ M, 5 μ M, 10 μ M). Fluorescence was analysed by flow cytometry.

Compound characterizations

Compound DBCO-ARM1



Figure S 4. Analytical RP-HPLC spectrum of compound DBCO-ABM1 (5-100% B in 15 minutes)



Figure S 5. Theoretical (top) and measured (bottom) HRMS spectrum of compound DBCO-ABM1



Figure S 6. ¹H NMR spectrum of compound **DBCO-ABM1** (D₂O, 500 MHz)



Compound DBCO-ABM4

Figure S 7. Analytical RP-HPLC spectrum of compound **DBCO-ABM4** (5-100% B in 15 minutes)



Figure S 8. Theoretical (top) and measured (bottom) HRMS spectrum of compound DBCO-ABM4



Figure S 9. ¹H NMR spectrum of compound **DBCO-ABM4** (D₂O, 500 MHz)



Compound DBCO-ABM16

Figure S 10. Analytical RP-HPLC spectrum of compound DBCO-ARM16 (0-80% B in 15 minutes)



Figure S 11. Theoretical (top) and measured (bottom) HRMS spectrum of compound DBCO-ARM16



Figure S 12. ¹H NMR spectrum of compound **DBCO-ABM16** (D₂O, 500 MHz)