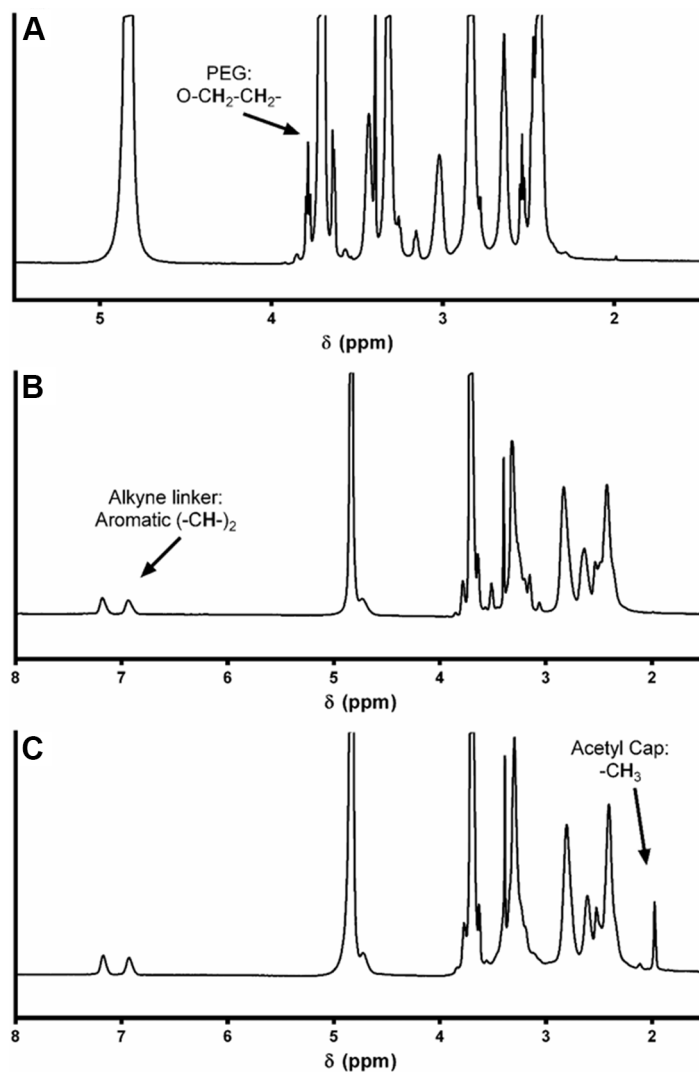
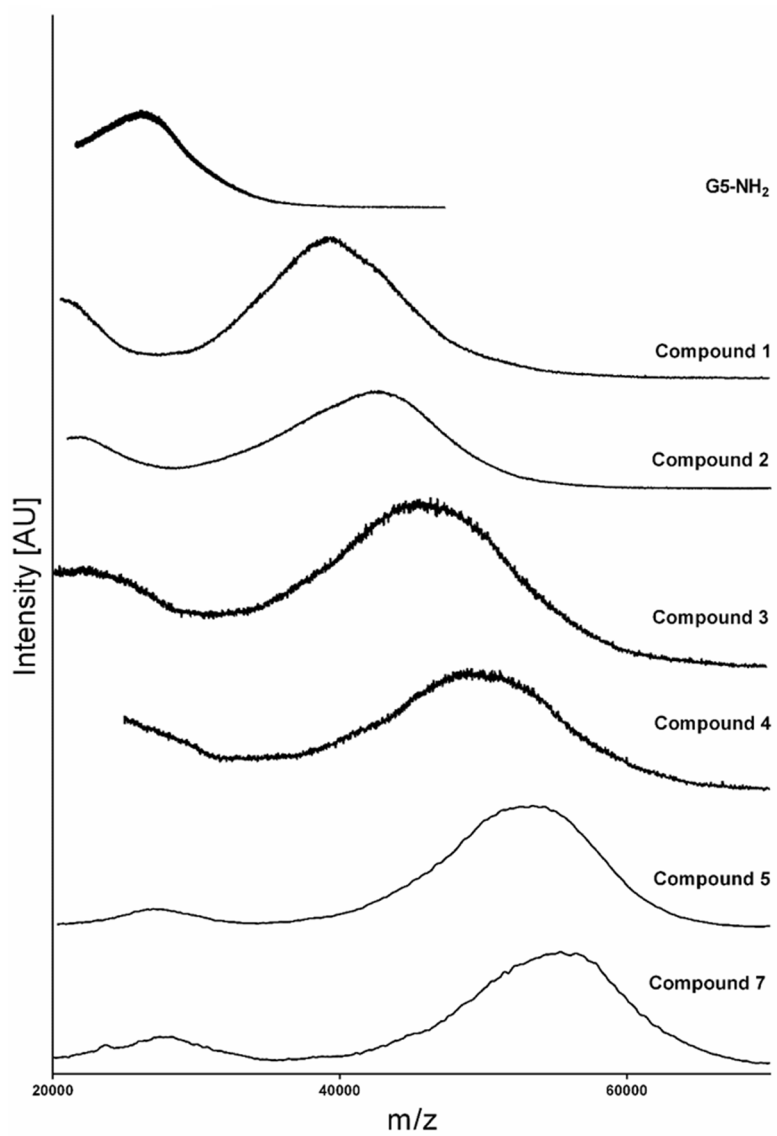


Dendrimer antibody conjugate to target and image HER-2 overexpressing cancer cells

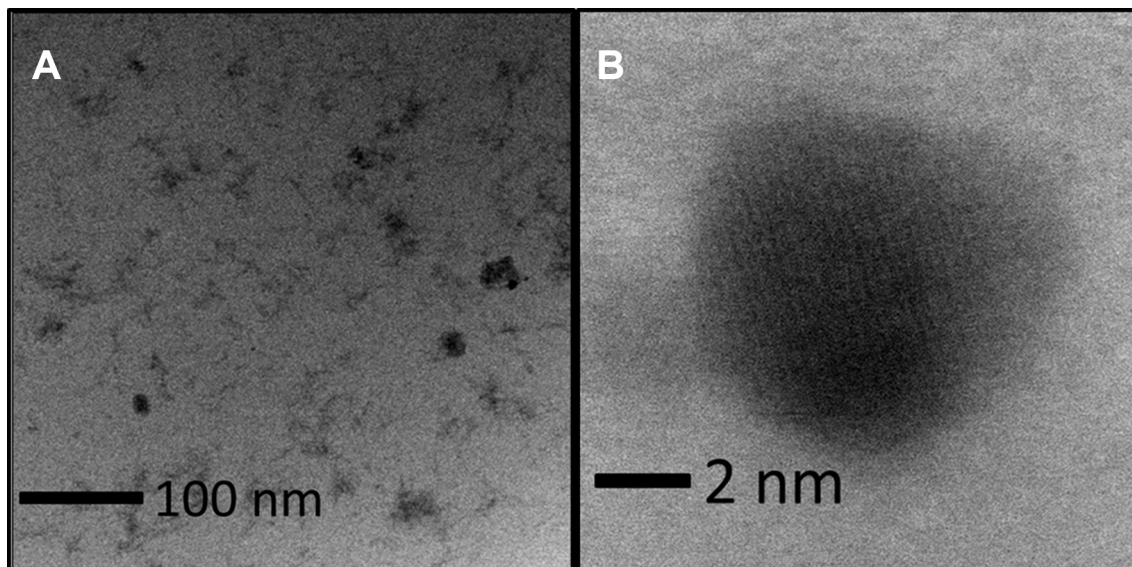
Supplementary Materials



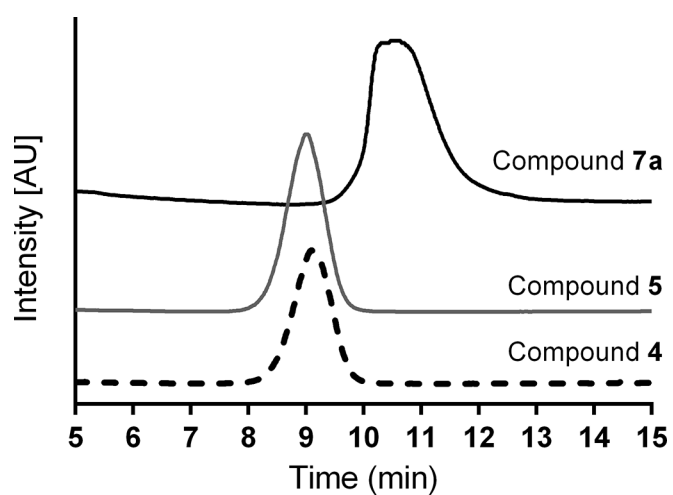
Supplementary Figure S1: NMR shifts for the initial dendrimer modification steps. (A) compound 1, (B) compound 2, (C) compound 4. Indicatory peaks of the respective modifications are labeled on each sub-figure. Samples were measured in 2 mg/mL D₂O.



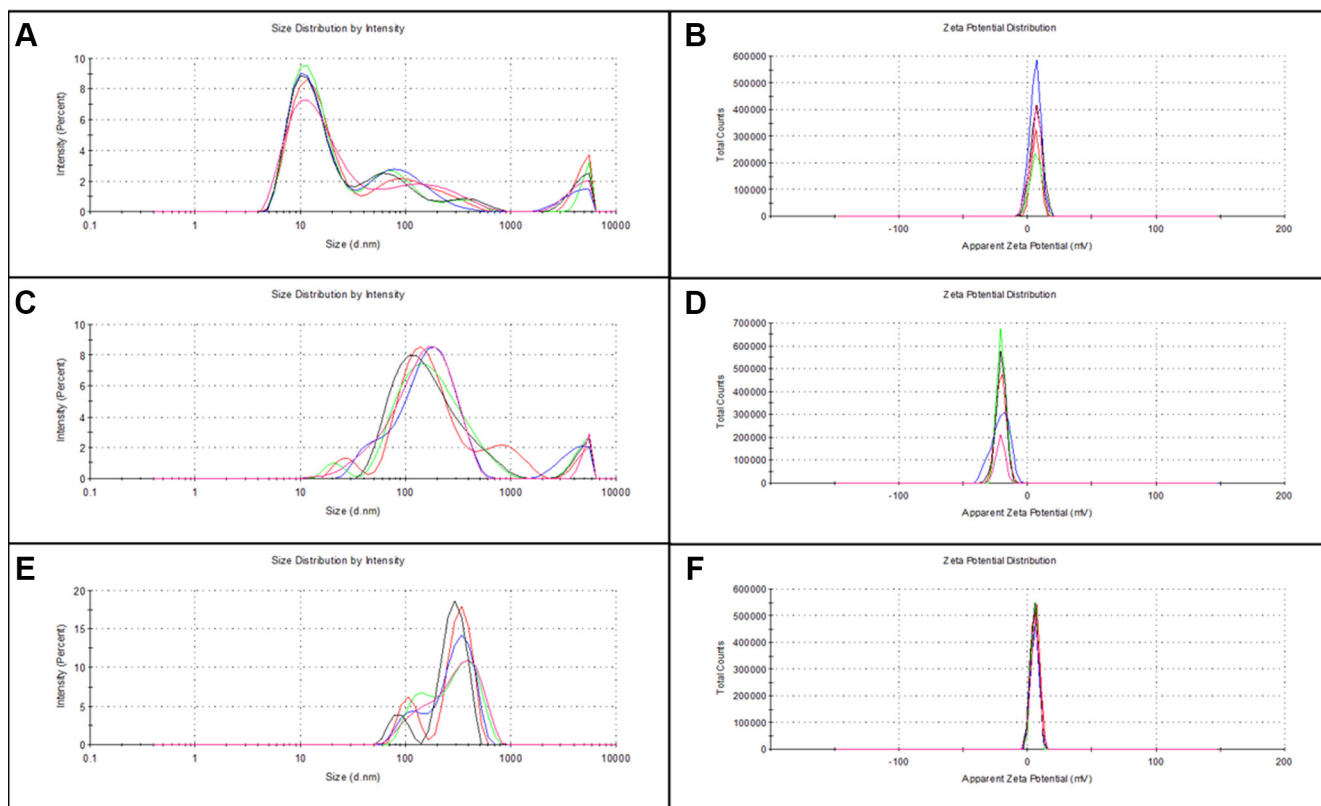
Supplementary Figure S2: Depicted are the MALDI-TOF spectra for each compound. The plot of 7 displays only the dendrimer's mass.



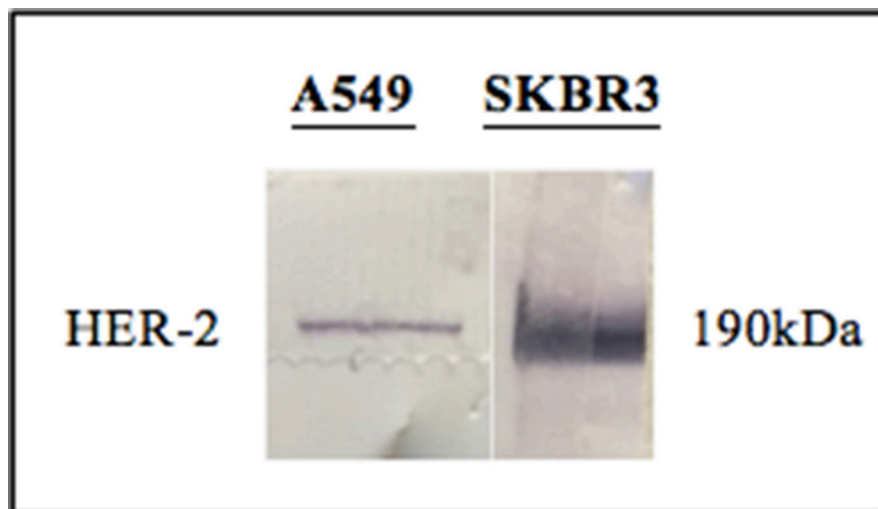
Supplementary Figure S3: TEM image of the dendrimer Herceptin conjugate. Image (A) depicts the dendrimer Herceptin complex with gadolinium and gold nanoparticles. Image (B) shows an AuNP at roughly five nanometers.



Supplementary Figure S4: UPLC Chromatograms of various compounds. (A) shows pure Herceptin, (B) shows 4, 5, and 7. Samples were dissolved in DI water and measured at 285 nm.



Supplementary Figure S5: (A) DLS plot of compound Au-G5-PEG-Alkyne-DOTA-Gd-NHAc (5) (28.3 ± 10.2 d. nm), (B) Zeta potential plot of compound 5 (6.3 ± 4.1 mV), (C) DLS plot of compound Herceptin-azide (6) (154.0 ± 7.0 d. nm), (D) Zeta potential plot of compound 6 (-20.86 ± 3.72 mV), E) DLS plot of Au-G5-Gd-Herceptin (7a) (459.0 ± 28 d. nm), F) Zeta potential plot of compound 7a (5.72 ± 3.0 mV). Samples were measured in 1mM HEPES buffer at 1mg/mL (reported as mean \pm one SD, $n = 5$).



Supplementary Figure S6: Western blot analysis of A549 and SKBR-3 cell lines. 15 μ g of total protein was loaded into each lane. The protein concentration was determined by BCA assay. The left lane: cell lysate of A549 cells; the right lane; cell lysate of SKBR-3 cells.