Supplementary Materials: Amino Acid-Modified Polyethylenimines with Enhanced Gene Delivery Efficiency and Biocompatibility

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Figure S1. Agarose gel retardation electrophoresis assay for release of DNA from polyplex in the presence of heparin at different weight ratio (heparin/DNA: 0, 0.25, 0.5, 1, 2, 4, 8, 16, 32; Ga-PEI/DNA: w/w = 3). The first lane is DNA control.



Figure S2. DNA protection against degradation by DNase. Lane 1: naked DNA; Lane 2: DNA treated with DNase for 2 h; Lane 3, 6, 9, 12, 15: polyplex (Polymer/DNA: w/w = 3); Lane 4, 7, 10, 13, 16: polyplex with DNase for 2 h at 37 °C; Lane 5, 8, 11, 14, 17: polyplex with DNase for 2 h at 37 °C followed by heat-inactivation of DNase and then treated with heparin.



Figure 3. Cont.



Figure S3. Particle size (**A**) and zeta potential (**B**) of polyplexes formed from Ga-PEI with different DS by DLS. Data represent mean \pm SD (n = 3).



Figure S4. Fluorescence microscope images of pEGFP-transfected U-2OS cells in the presence of 10% serum at optimal weight ratio.



Figure S5. Fluorescence microscope images of pEGFP-transfected 293 cells in the absence of serum at optimal weight ratio.



Figure S6. Fluorescence microscope images of pEGFP-transfected 293 cells in the presence of 10% serum at optimal weight ratio.



Figure S7. Fluorescence microscope images of pEGFP-transfected HeLa cells in the absence of serum at optimal weight ratio.



Figure S8. Fluorescence microscope image of pEGFP-transfected HeLa cells in the presence of 10% serum at optimal weight ratio.



Figure S9. Cytotoxicity of polycations at different concentration toward U-2OS cells: effect of DS. Data represent mean \pm SD (*n* = 3).

¹H NMR Spectra:























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