Electronic Supporting Information

Formation of iron oxide nanoparticle-loaded γ-polyglutamic acid nanogels for MR imaging of tumors[†]

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Fig. S1. (a) Hydrodynamic size distribution of the γ -PGA/PEI-Fe₃O₄ NGs in water; (b) hydrodynamic size of the γ -PGA/PEI-Fe₃O₄ NGs in water as a function of storage time.



Fig. S2. Hydrodynamic size of the γ -PGA/PEI-Fe₃O₄ NGs in acidic aqueous solution (pH = 6.5) and in acidic aqueous solution (pH = 6.5) containing high GSH concentrations (10 or 30 μ M), respectively.



Fig. S3. XRD pattern of γ -PGA/PEI-Fe₃O₄ NGs.



Fig. S4. TGA curves of naked Fe₃O₄ NPs, PEI-Fe₃O₄ NPs, and γ-PGA/PEI-Fe₃O₄ NGs.



Fig. S5. Phase contrast microscopic images of HeLa cells treated with PBS (a), and γ -PGA/PEI-Fe₃O₄ NGs at Fe concentrations of 0.01 mM (b), 0.02 mM (c), 0.04 mM (d), 0.06 mM (e), 0.08 mM (f), 0.1 mM (g), and 0.2 mM (h) for 24 h.



Fig. S6. (a) Biodistribution of Fe in the major organs of the nude mice including the heart, liver, spleen, lung, kidney, and tumor at 2, 4, 6, 12, and 24 h post-intravenous injection of the γ -PGA/PEI-Fe₃O₄ NGs ([Fe] = 51.04 mM, 0.2 mL PBS, for each mouse); (b) Biodistribution of Fe in the major organs of the healthy mice including the heart, liver, spleen, lung, and kidney at 7, 14, and 30 days post-intravenous injection of the γ -PGA/PEI-Fe₃O₄ NGs ([Fe] = 51.04 mM, 0.2 mL PBS, for each mouse); (b) Biodistribution of Fe in the major organs of the healthy mice including the heart, liver, spleen, lung, and kidney at 7, 14, and 30 days post-intravenous injection of the γ -PGA/PEI-Fe₃O₄ NGs ([Fe] = 51.04 mM, 0.2 mL PBS, for each mouse).