

## Supporting information

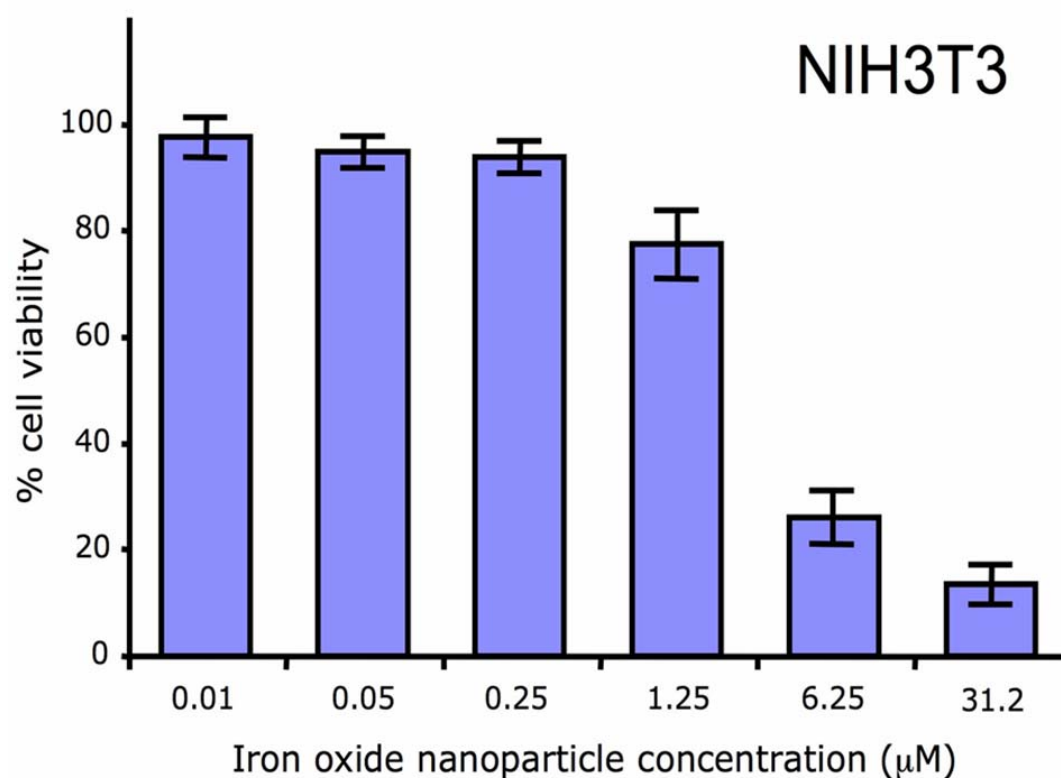
### **Modularly Assembled Magnetite Nanoparticles Enhance in vivo Targeting for Magnetic Resonance Cancer Imaging**

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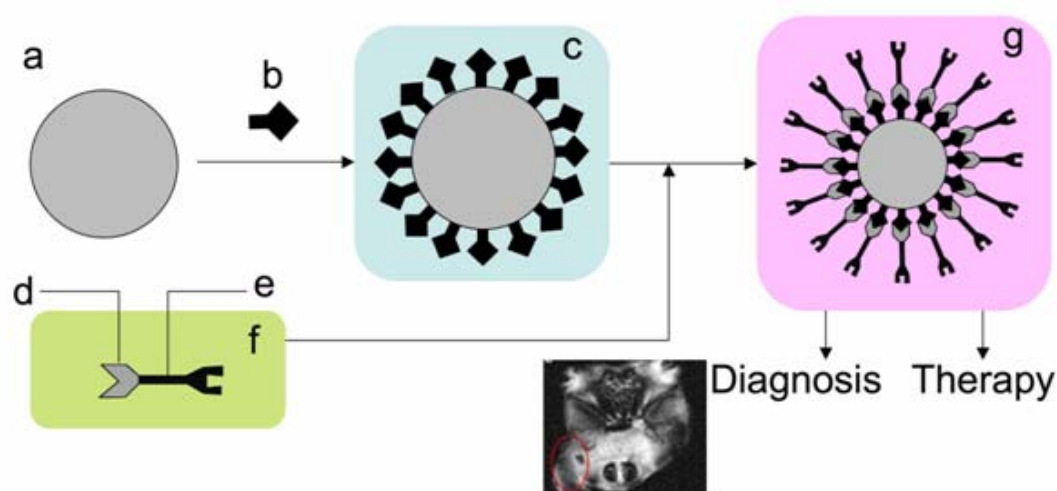
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S1.....Figure S1

S2.....Figure S2



**Figure S1.** To evaluate the biocompatibility of the nanoparticles, NIH3T3 cells were cultured in 96-well plates. Fe<sub>3</sub>O<sub>4</sub>-NTA-Ni-RGD4C nanoparticles at different concentrations (range: 0.01-31 μM) were added to each culture well and incubated for an additional 24 h before the spectrophotometric quantification of cell viability using an MTT assay. Cell viability was satisfactory (> 80%) at a particle concentration below 1.25 μM.



**Figure S2.** We developed a new concept of modular-targeting functional nanoparticles through self-assembly of the targeting moiety and the functional nanoparticles by modifying the nanoparticle surface with a small "key molecule" and the targeting peptide or recombinant antibody with an affinity peptide tag for the "lock molecule".

Schematic diagram illustrating the concept of modular-targeting functional nanoparticles for diagnostic imaging and therapeutic applications. Nanoparticle module (c) is composed of a functional nanoparticle (a) with a common surface "key molecule" (b), Ni-NTA for example, to self-assemble with targeting moieties in biomolecular module (d~f). Biomolecular module (f) is composed of a biomolecule (d) such as antibody or a peptide with a genetically engineered "lock molecule" (e) tag (6-his domain for example) for binding to "key molecules" on the nanoparticles. The two modules self-assembled to form functional targeting nanoparticles (g) that not only gives a versatile combination for biological end and the nanoparticle end based on clinical applications , but also may improve the targeting because of the orientation-control and enrichment of the effective targeting moiety through self-assembly.