

Supporting Information

Superparamagnetic Hollow and Paramagnetic Porous Gd_2O_3 Particles

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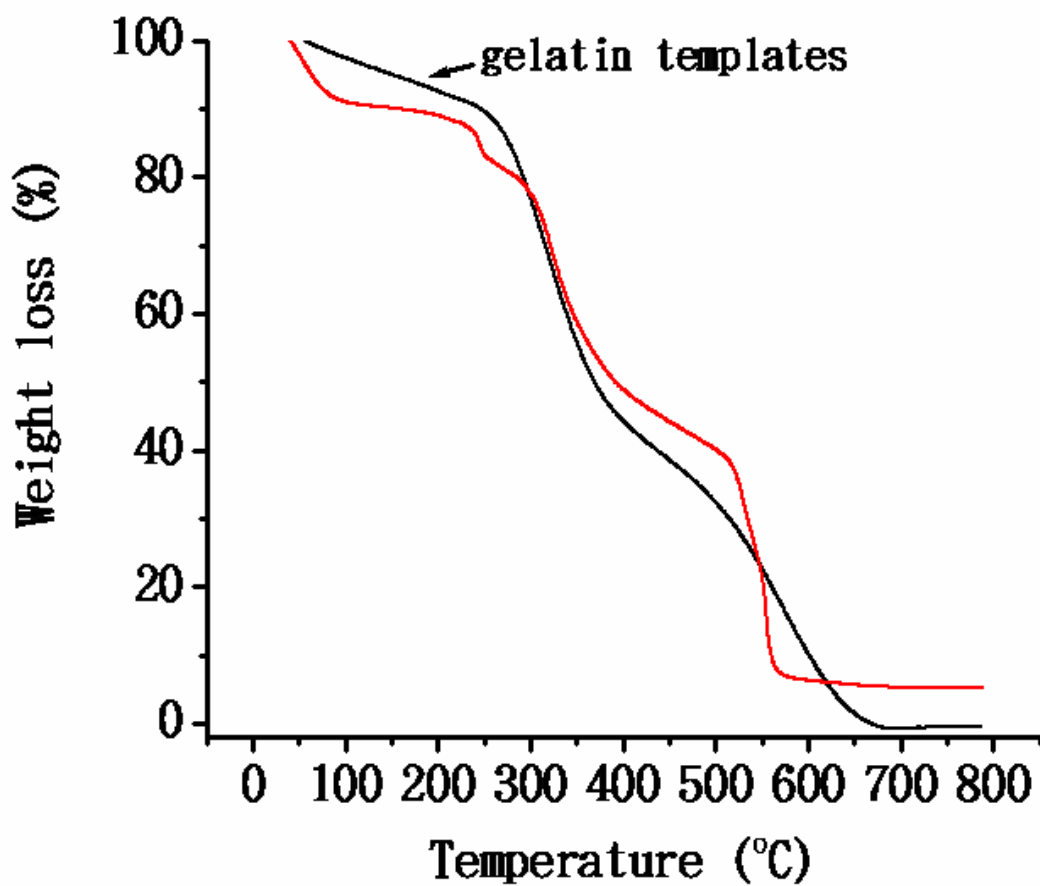


Figure S1. TGA measurement of as-synthesized gelatin templates (black line) and the gelatin/inorganic composites (red line). The gelatin/inorganic composites were prepared through method 1 (sol-gel process) by hydrolysis of organometallic gadolinium isopropoxide over the gelatin cores under hydrothermal conditions.

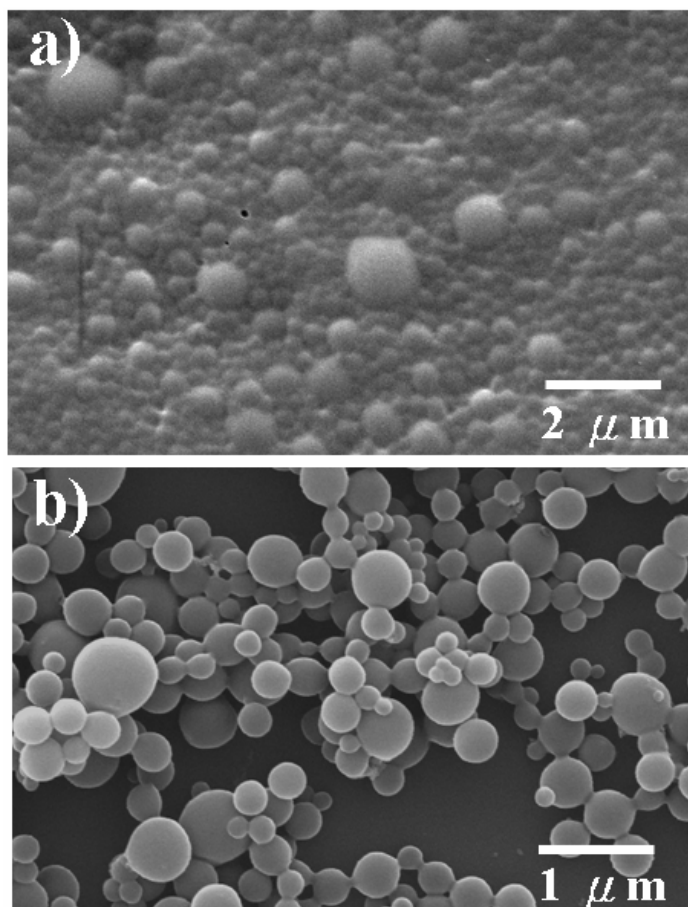


Figure S2. SEM images of the as-synthesized gelatin templates a) and the gelatin/inorganic composites b). The gelatin/inorganic composites were prepared using method 1 (sol-gel process) by hydrolyzing organometallic gadolinium isopropoxide over the gelatin cores under hydrothermal conditions.

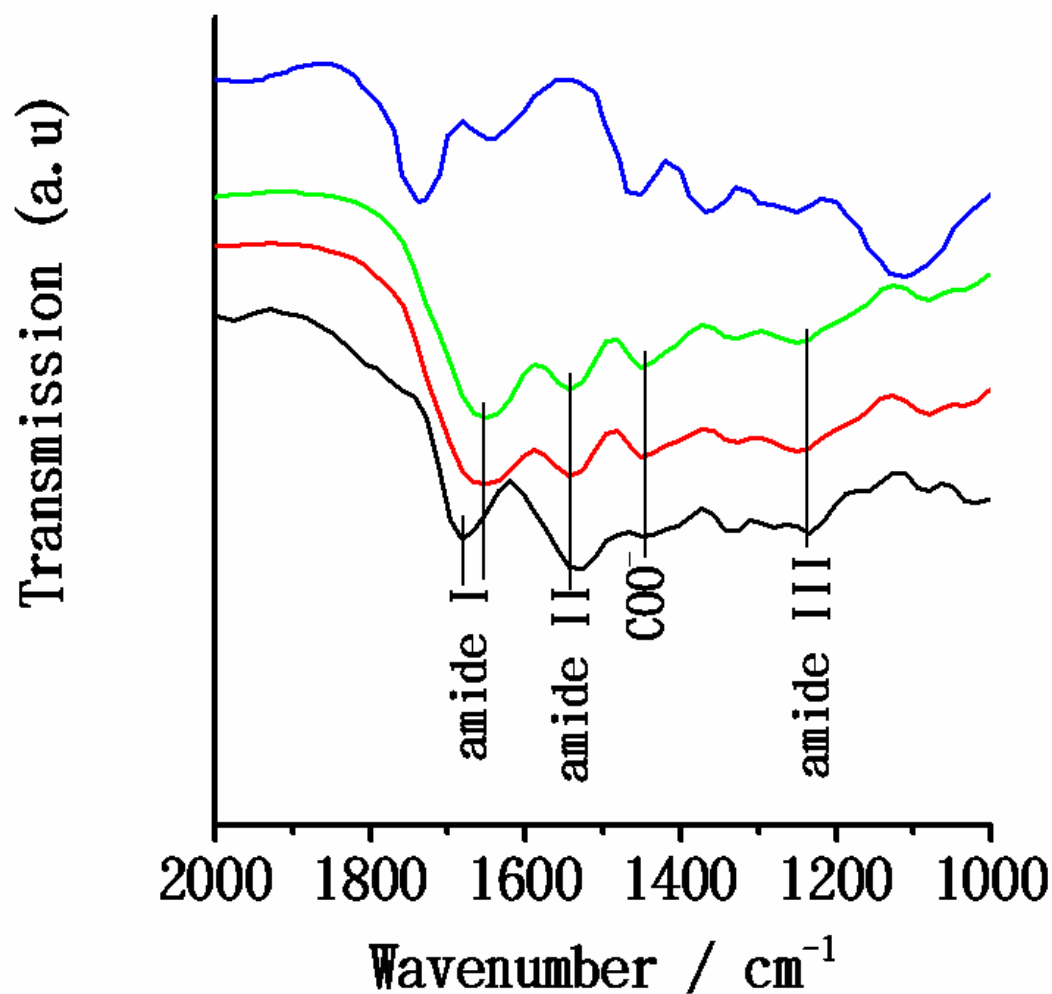


Figure S3. IR spectrum of gelatin powder (black line), as-synthesized gelatin templates (red line), inorganic/gelatin composites (green line), and P123 polymer (blue line). The gelatin powder purchased from Acros was used as a standard with which to compare the as-synthesized gelatin templates. The gelatin/inorganic composites were prepared using method 1 (sol-gel process) by hydrolyzing organometallic gadolinium isopropoxide over the gelatin cores under hydrothermal conditions.

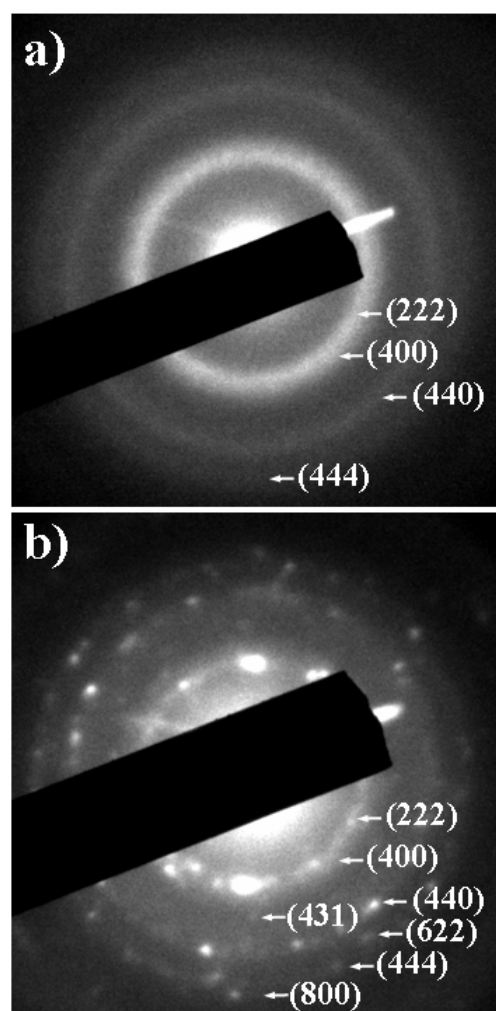


Figure S4. Selected area electron diffraction patterns (SAED) of hollow a) and porous b) Gd_2O_3 nanospheres.

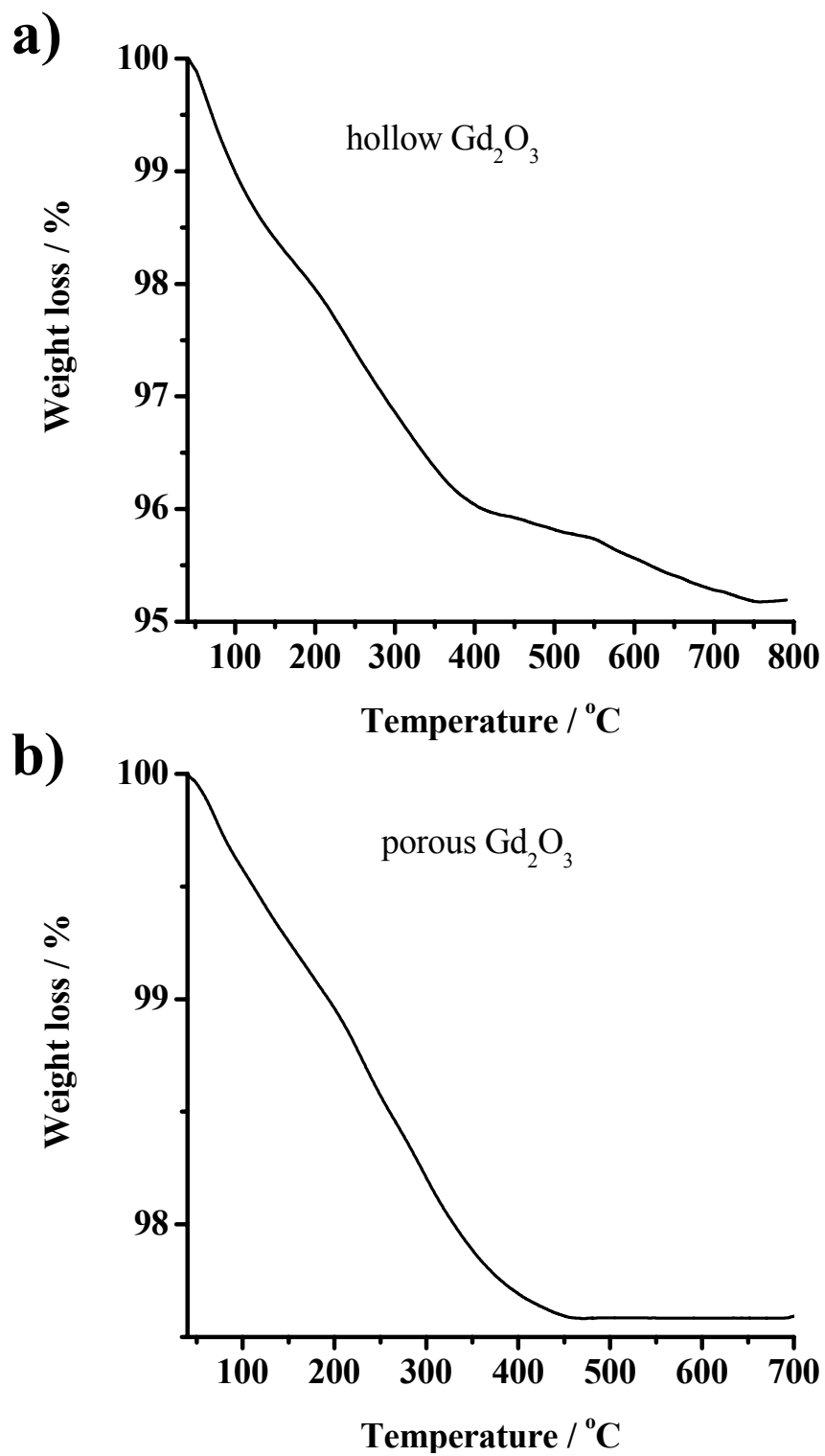


Figure S5. TGA measurements of the hollow (a) and porous (b) Gd_2O_3 nanospheres.

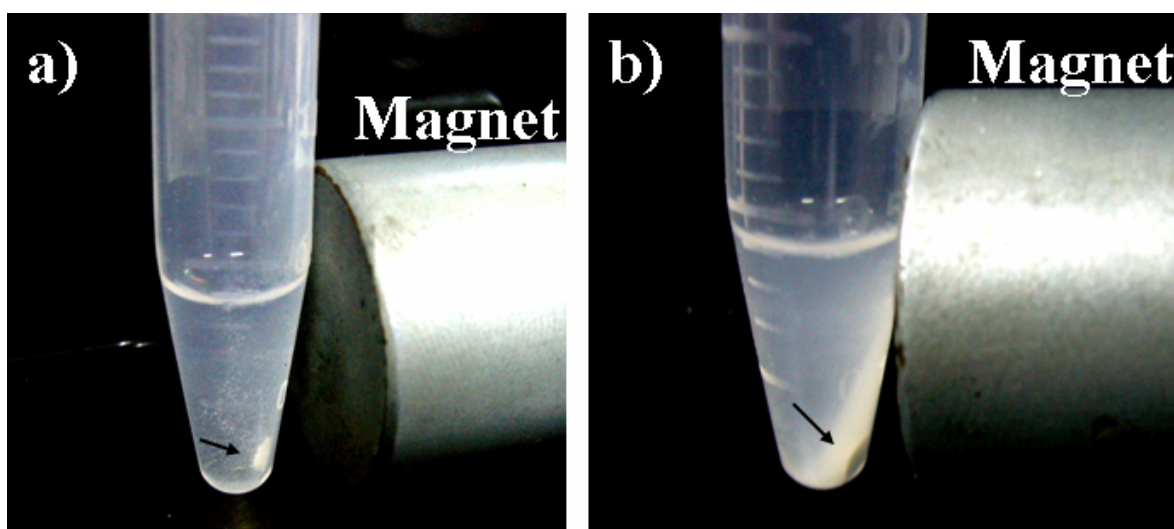


Figure S6. The attraction of a) hollow Gd_2O_3 nanospheres and b) porous Gd_2O_3 nanospheres under an external magnetic field.

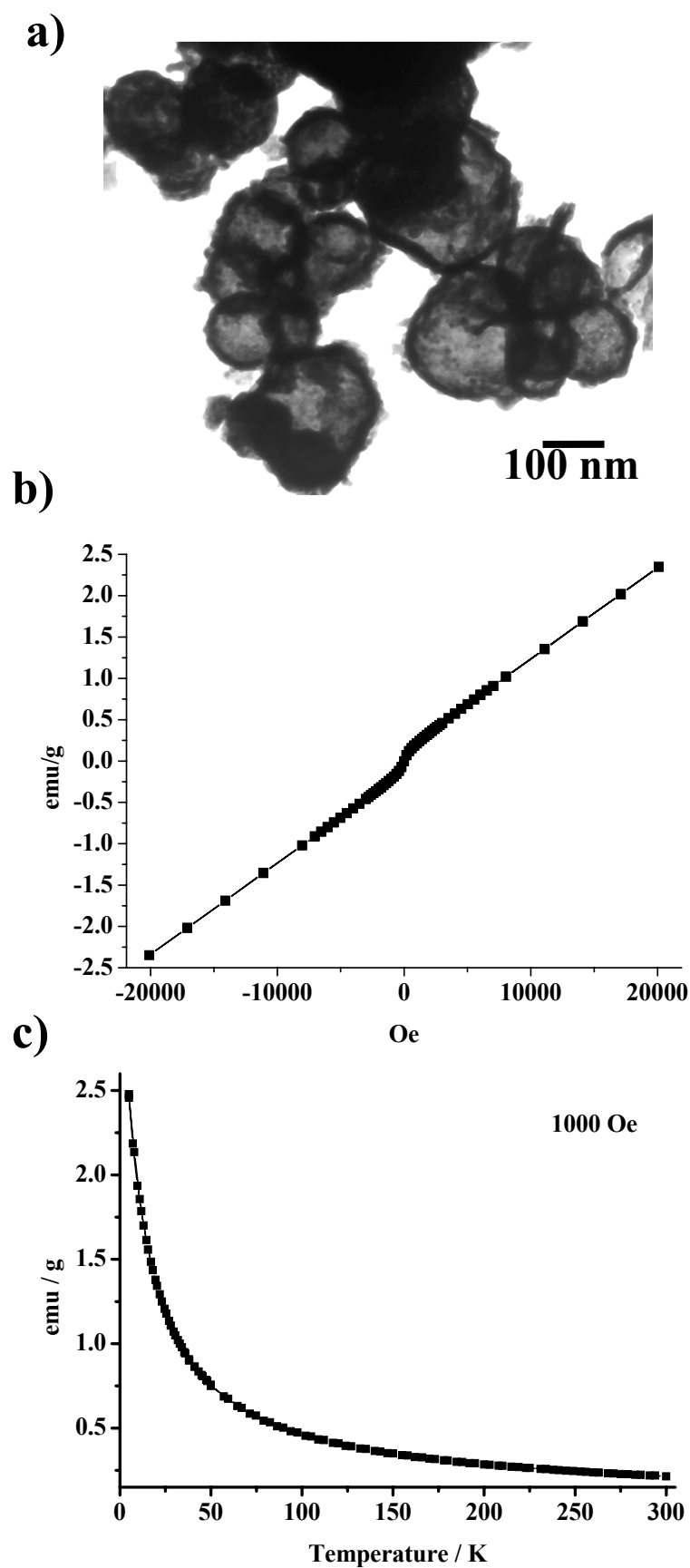
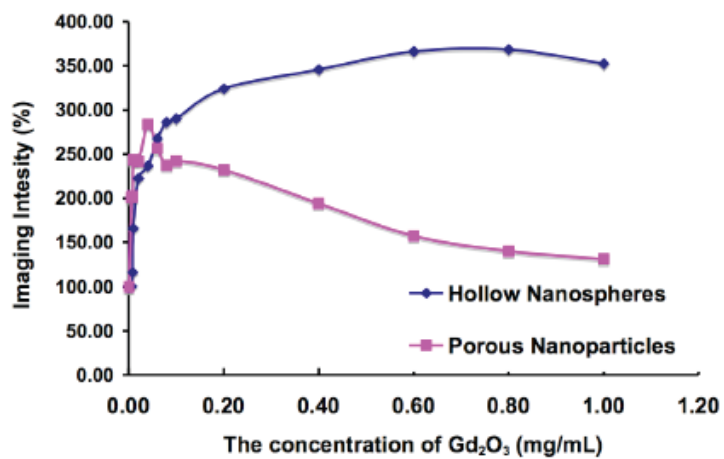


Figure S7. (a) TEM image, (b) Magnetization ($M-H$) plot at 300 K, and (c) ZFC-FC ($M-T$) curves at 1000 Oe of calcined hollow Gd_2O_3 nanosphere (800 °C).

a)



b)

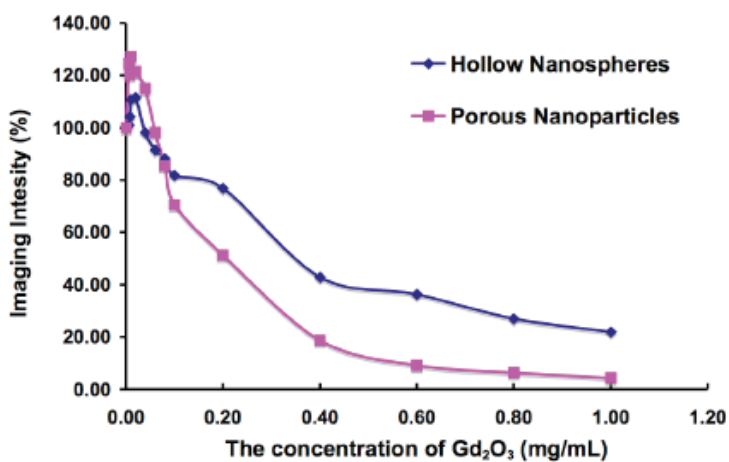


Figure S8. The signal intensity of a) T₁- and b) T₂-weighted imaging for hollow and porous Gd₂O₃ nanospheres.

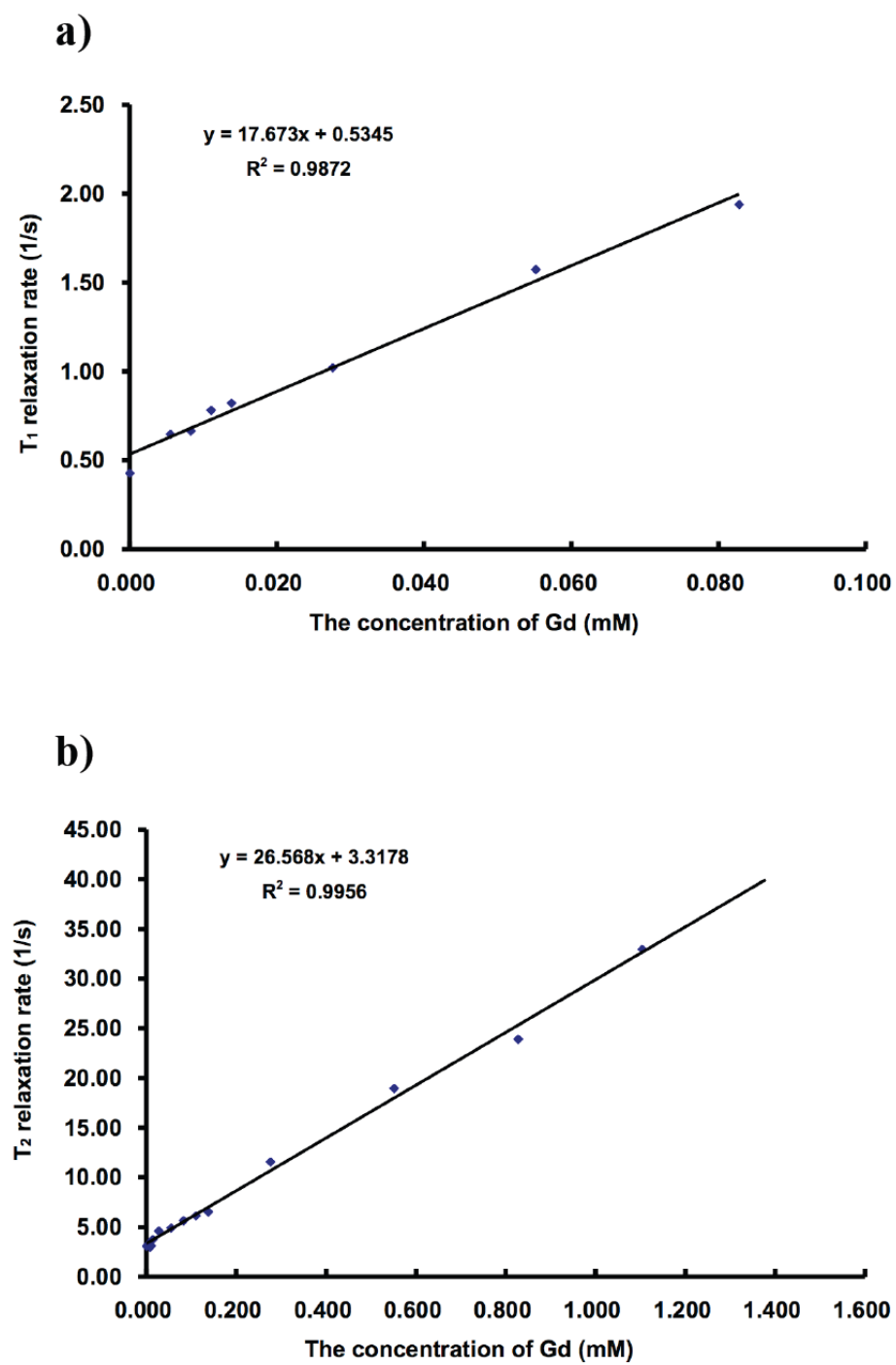


Figure S9. The r_1 a) and r_2 b) relaxivity curves obtained using various concentrations of hollow Gd₂O₃ nanospheres in 5% agarose gel.

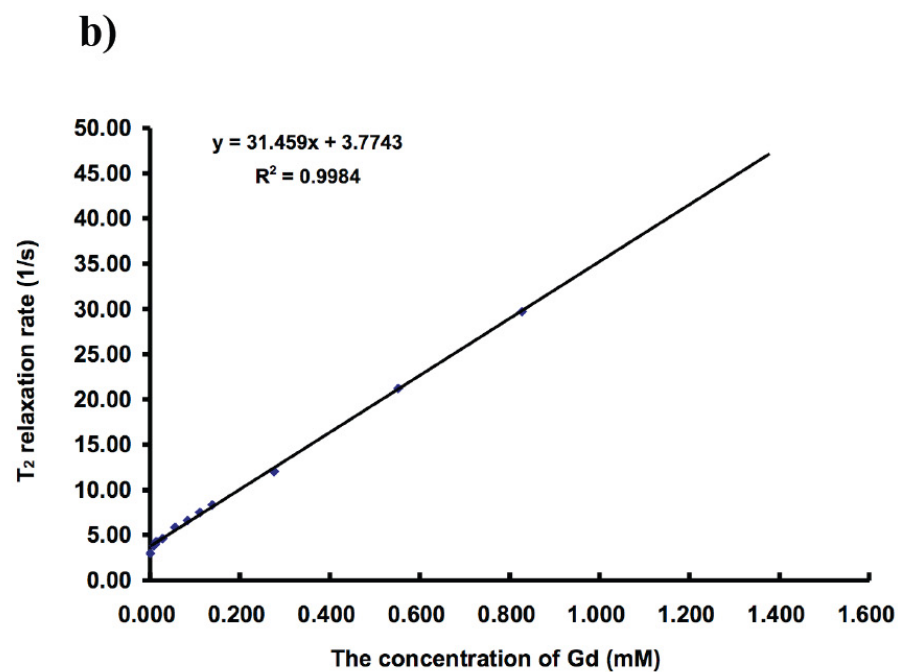
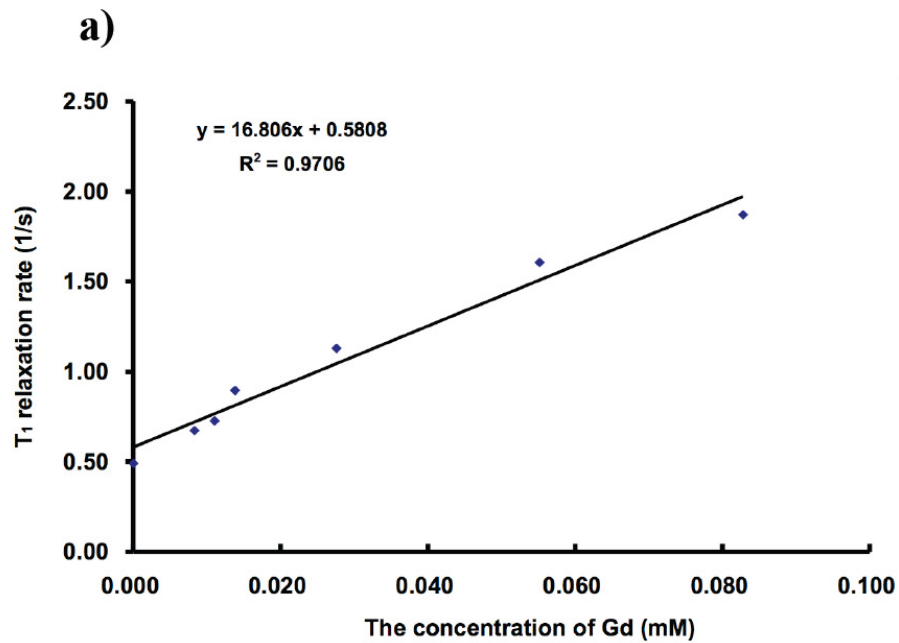


Figure S10. The r_1 a) and r_2 b) relaxivity curves obtained using various concentrations of porous Gd_2O_3 nanospheres in 5% agarose gel.

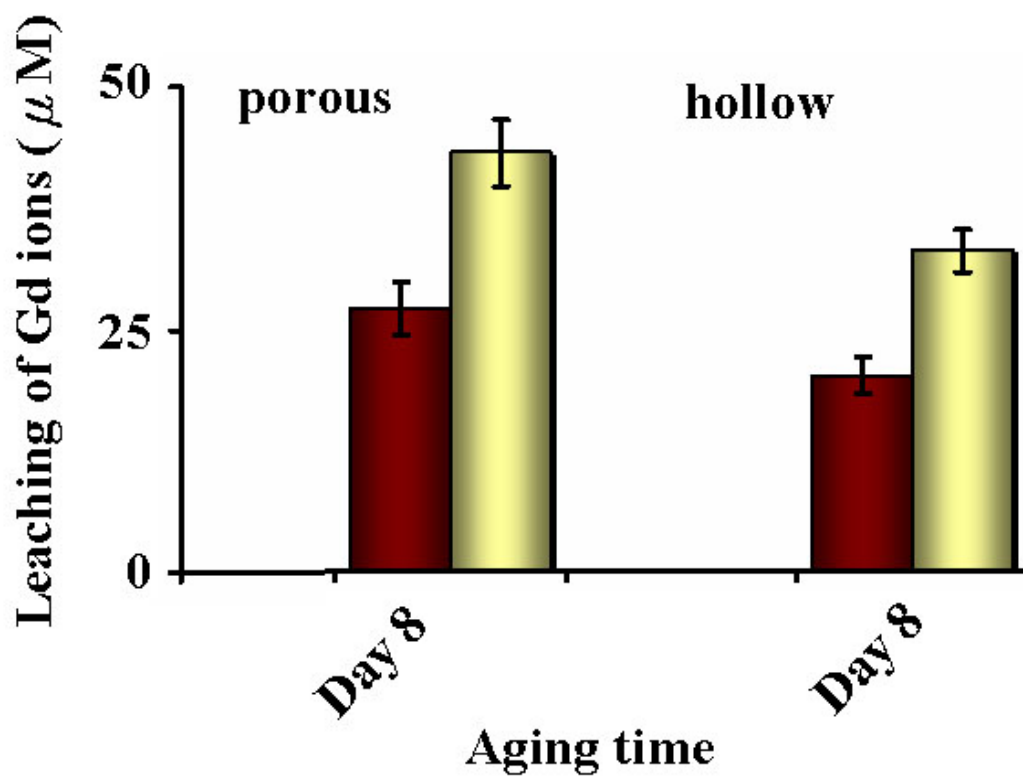


Figure S11. Leaching Gd³⁺ ions from hollow (right) and porous (left) Gd₂O₃ nanospheres, which was analyzed by ICP-AES measurements after the nanospheres had been stored at 4 °C (red column) and 37 °C (yellow column) for 8 days.

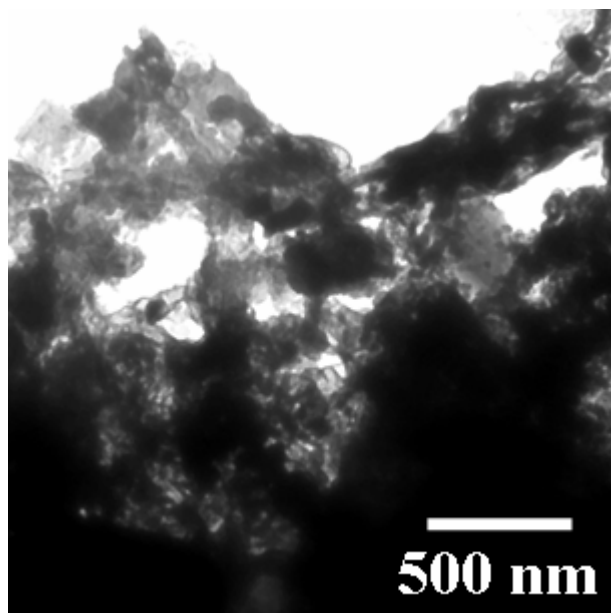


Figure S12. The TEM image shows the result of preparing hollow Gd_2O_3 nanospheres using method 1 (sol-gel process) but without hydrothermal treatment.

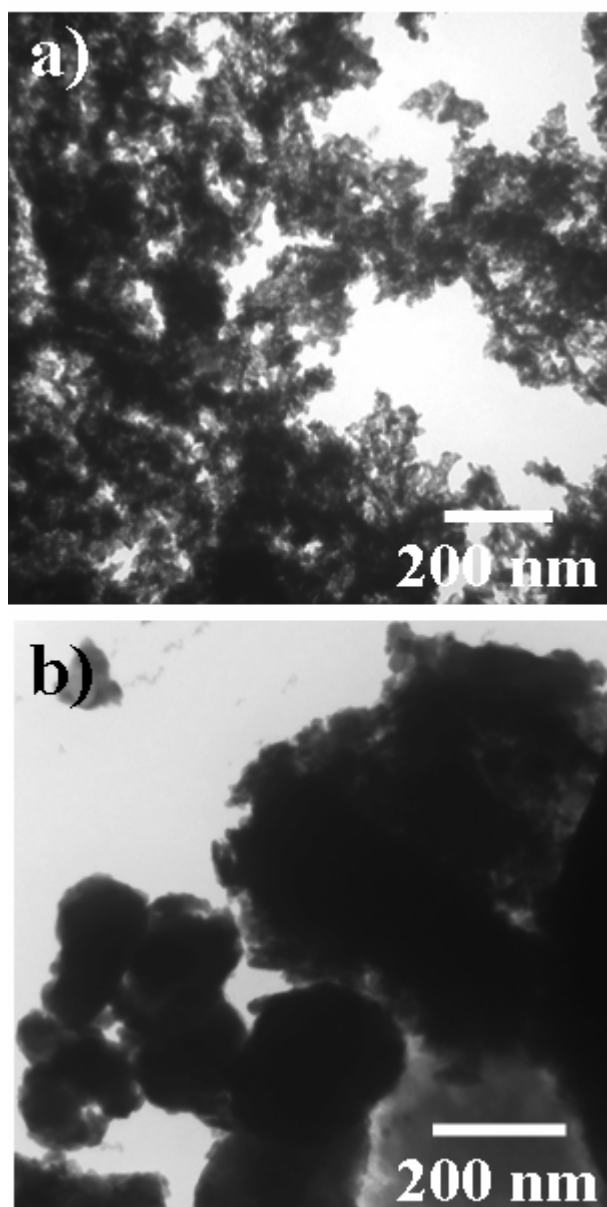


Figure S13. TEM images of Gd₂O₃ nanospheres prepared using a) method 1 (sol-gel process) and b) method 2 (precursor deposition) but without P123 polymer.

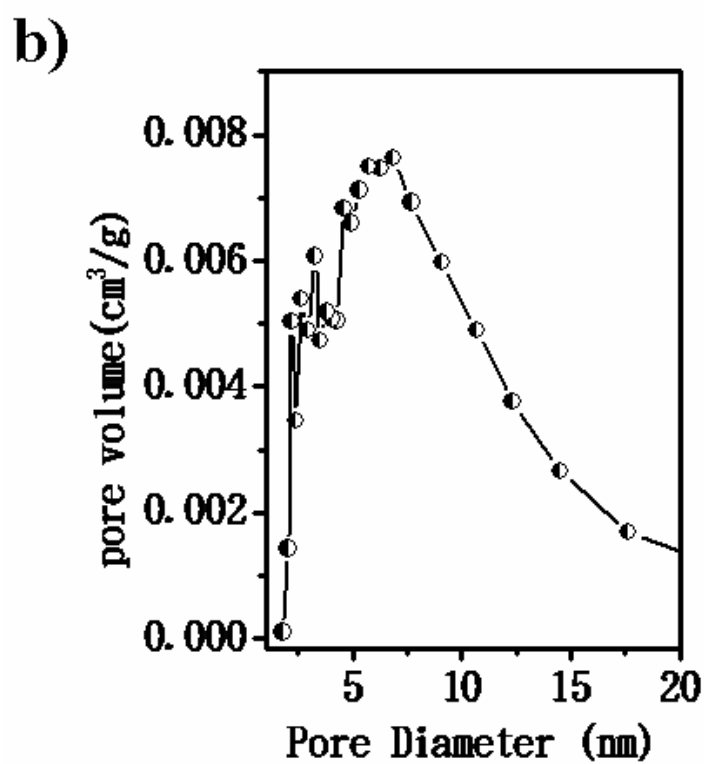
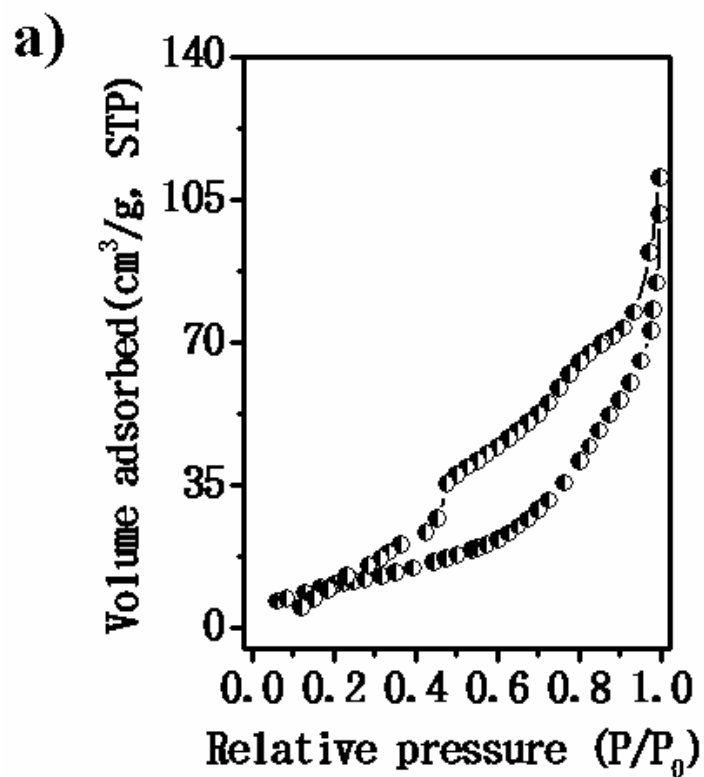


Figure S14. a) N_2 adsorption-desorption isotherms and b) BJH pore size distributions of porous TiO_2 nanospheres.

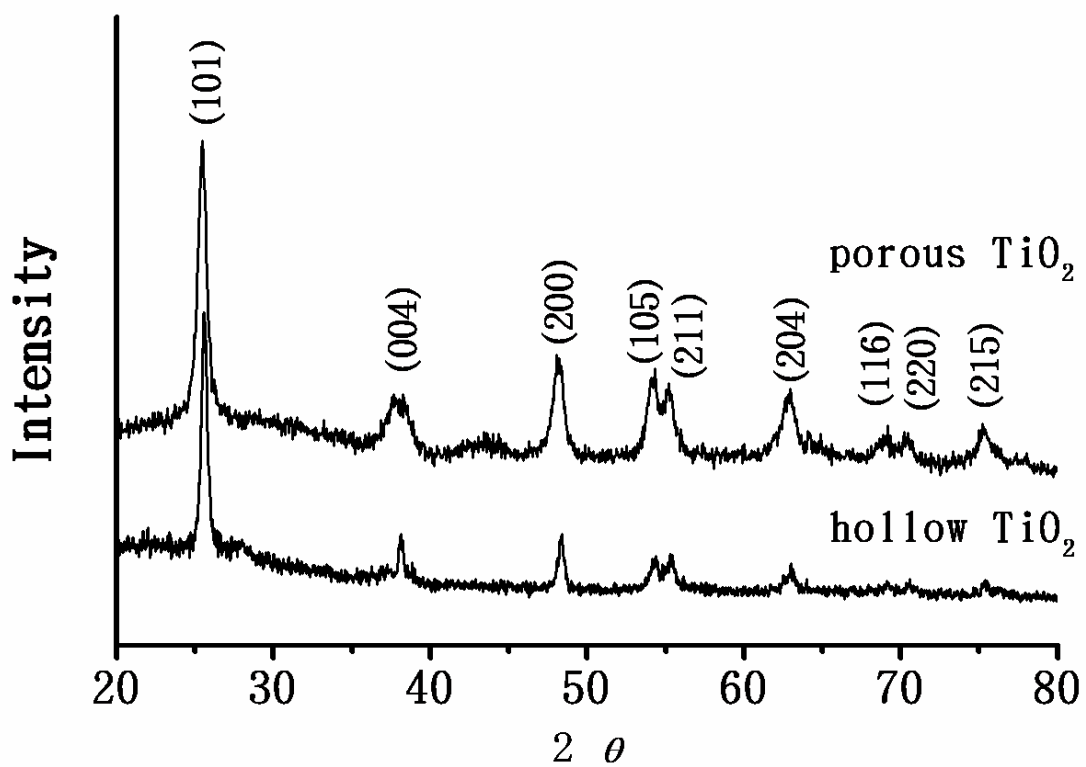


Figure S15. XRD patterns of hollow and porous TiO₂ nanospheres.